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(54) **DATA COMMUNICATION AND CONTROL FOR MEDICAL IMAGING SYSTEMS**

(75) Inventors: **Arthur E. Uber, III**, Pittsburgh;
Edward J. Rhinehart, Monroeville;
Rosemary Almon-Martin, Saxonburg;
David M. Griffiths, Pittsburgh; **Alan D. Hirschman**, Glenshaw; **Francis J. Sciulli**, Crafton; **Walter Grumski**, Pittsburgh; **Vera Pagano**, Bellevue; **Karen Zelenski**, Seven Fields, all of PA (US)

(73) Assignee: **Medrad, Inc.**, Indianola, PA (US)

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(52) **U.S. Cl.** **600/431; 604/31; 604/67**

(58) **Field of Search** 600/420, 431, 600/432, 458; 604/20, 31, 65-67, 246

(56) **References Cited**

U.S. PATENT DOCUMENTS

4,477,923 A 10/1984 Baumann et al.
4,854,324 A * 8/1989 Hirschman et al. 600/432
4,913,154 A * 4/1990 Ermert et al. 600/431

4,922,916 A * 5/1990 Ermert et al. 600/420
5,301,672 A * 4/1994 Kalender 600/431
5,417,213 A * 5/1995 Prince 600/420
5,485,831 A * 1/1996 Holdsworth et al. 600/431
5,544,215 A 8/1996 Shroy, Jr. et al.
5,799,649 A * 9/1998 Prince 600/420
5,827,504 A 10/1998 Yan et al.
5,885,216 A * 3/1999 Evans, III et al. 600/431

FOREIGN PATENT DOCUMENTS

DE 32 03 594 A1 8/1983
DE 197 02 896 A1 7/1997
DE 196 47 701 A1 5/1998
GB 2 328 745 A 3/1999

* cited by examiner

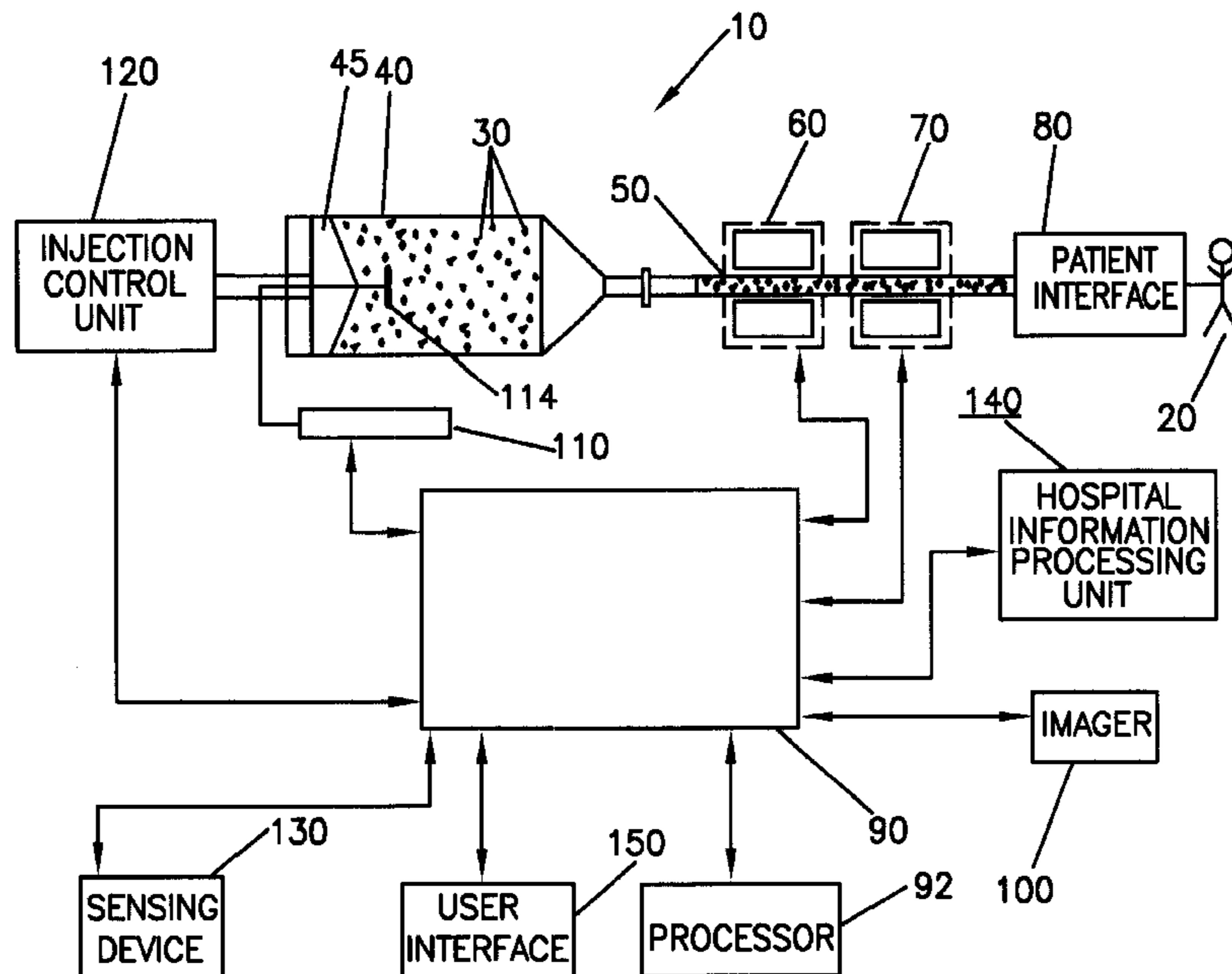
Primary Examiner—Brian L. Casler

(74) *Attorney, Agent, or Firm*—Henry E. Bartony, Jr.; Gregory L. Bradley

(57) **ABSTRACT**

A system for producing a contrast-enhanced medical image of a patient includes a source of a contrast or enhancement medium, a pressurizing unit in fluid connection with the source of contrast or enhancement medium, an energy source operable to apply energy to a region of the patient, an imaging unit providing a visual display of an internal view of the patient based upon a signal resulting from the energy applied to the region of the patient, and a control unit. In an embodiment, the signal is affected by a condition of the contrast or enhancement medium in the patient. To control the procedures, the control unit adjusts the condition of the contrast or enhancement medium in the patient based upon the signal. A communication interface preferably enables information between an injector subsystem and an imaging subsystem.

53 Claims, 6 Drawing Sheets



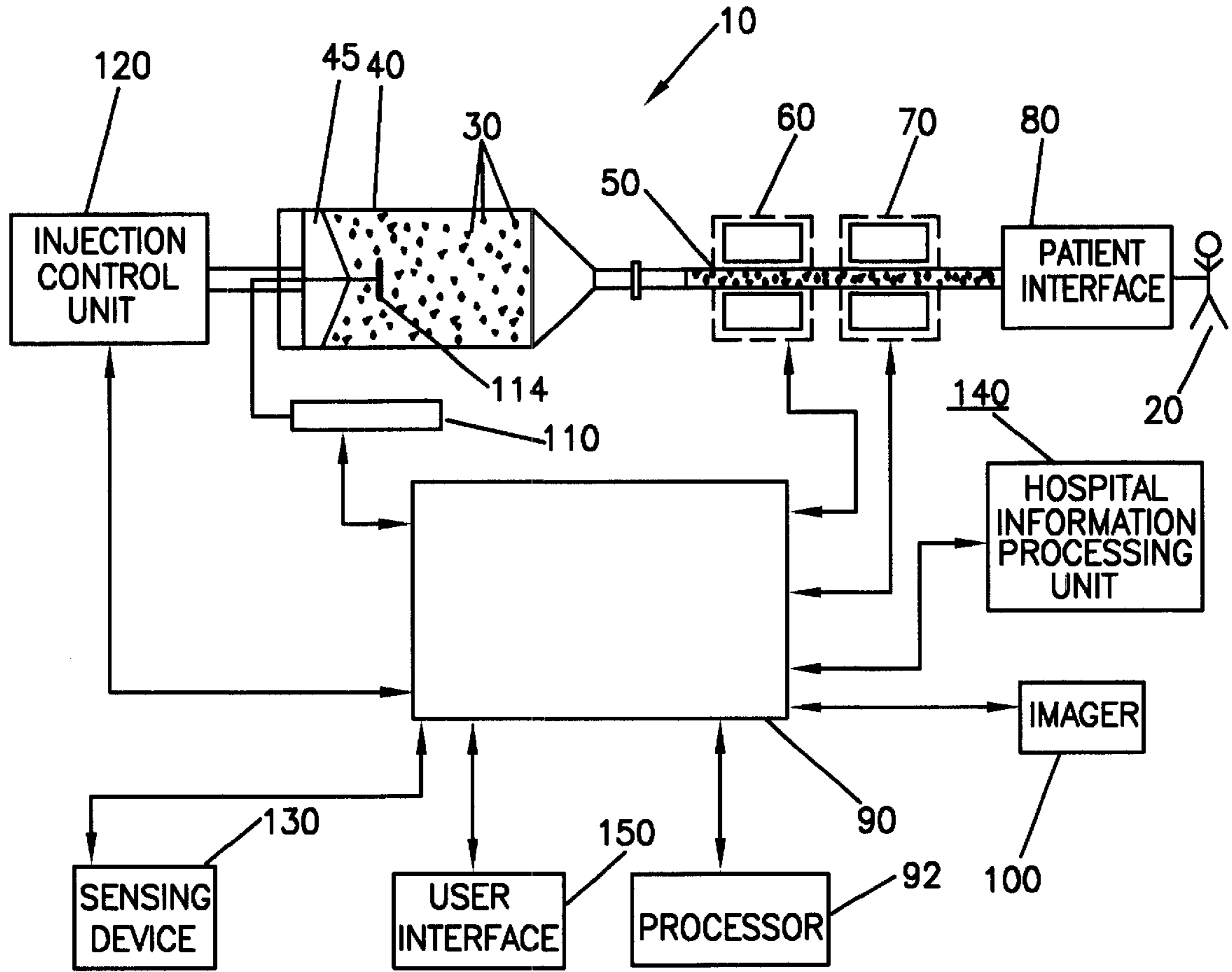


FIG. 1

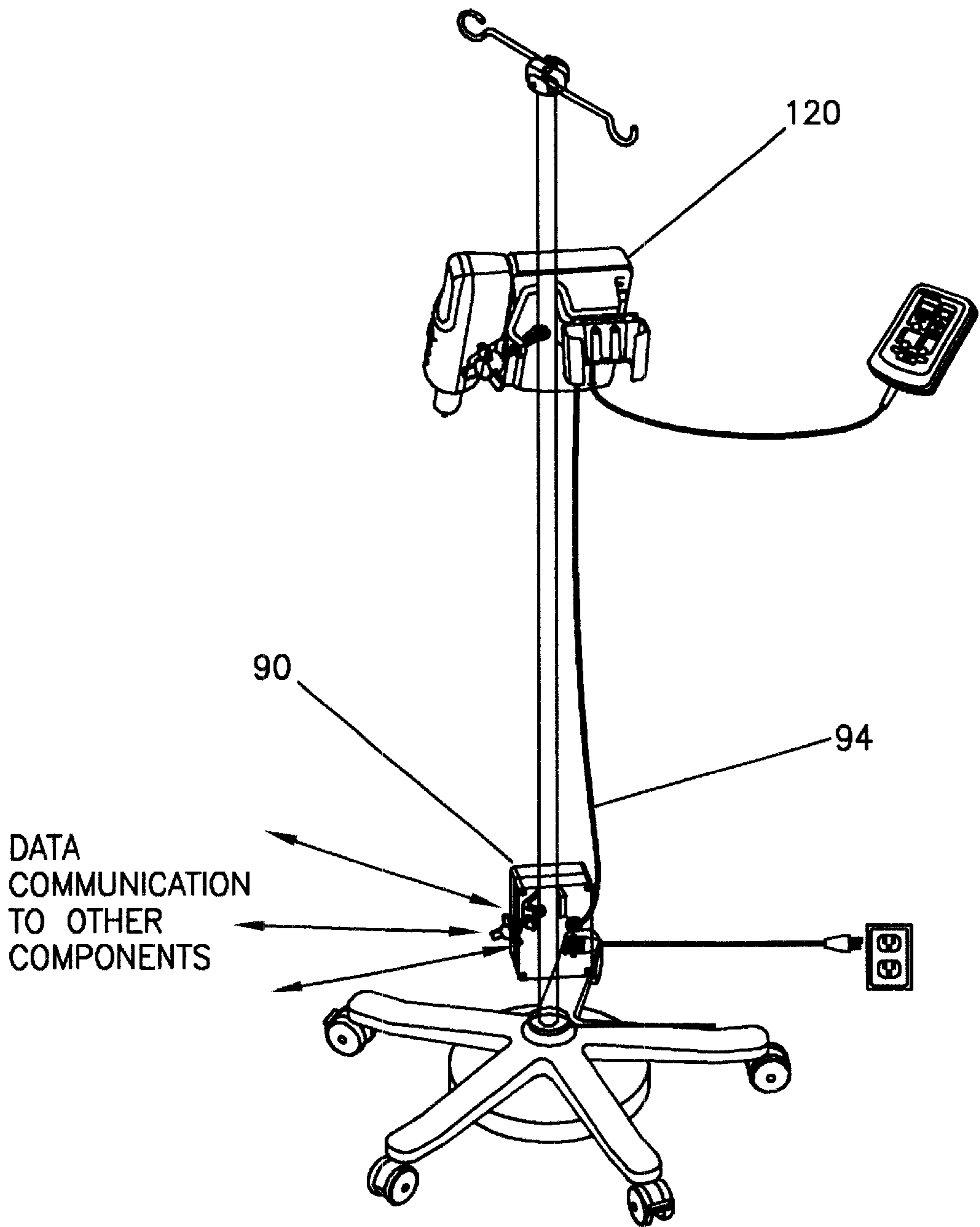
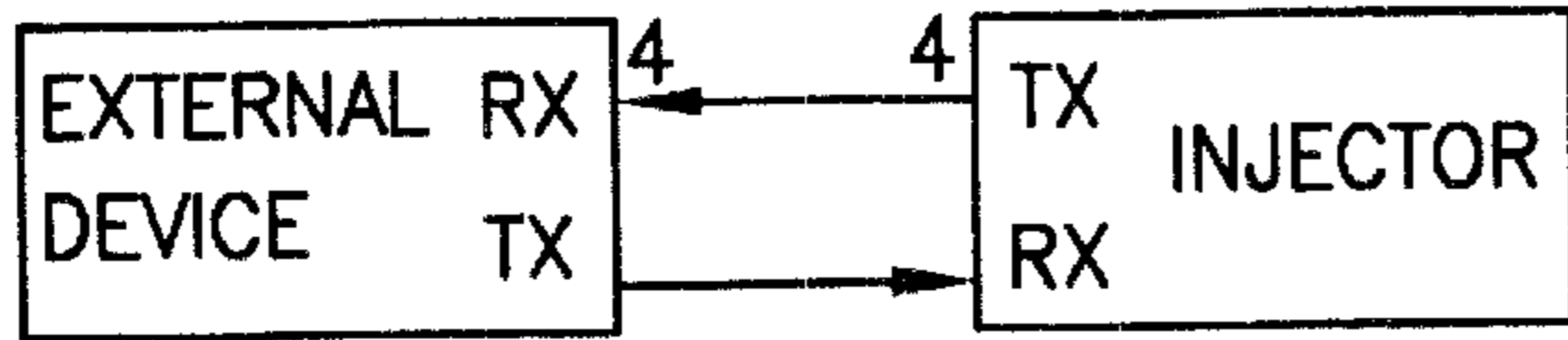
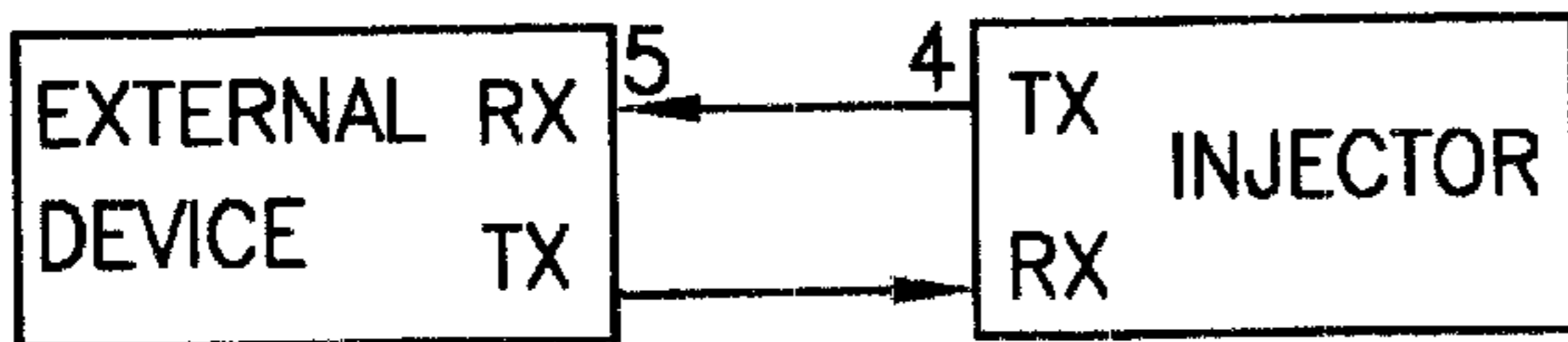


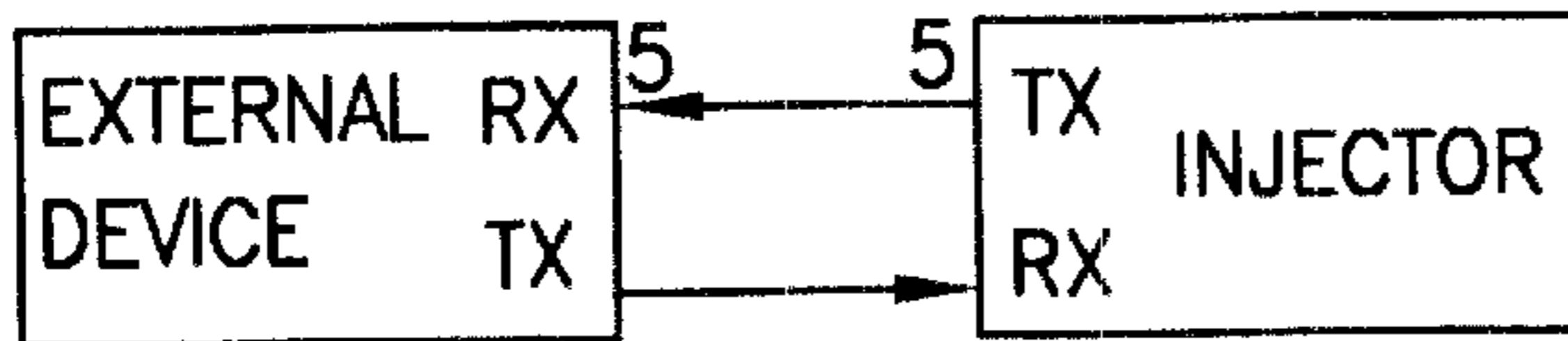
FIG. 2



SEQUENCE NUMBERS BEFORE MESSAGE TRANSMISSION.



SEQUENCE NUMBERS AFTER THE INJECTOR HAS SENT A MESSAGE AND THE EXTERNAL DEVICE RECEIVED AND VALIDATED IT.



SEQUENCE NUMBERS AFTER EXTERNAL DEVICE ACKNOWLEDGED THE RECEIVED MESSAGE AND THE INJECTOR RECEIVED THE ACKNOWLEDGE.

FIG. 4

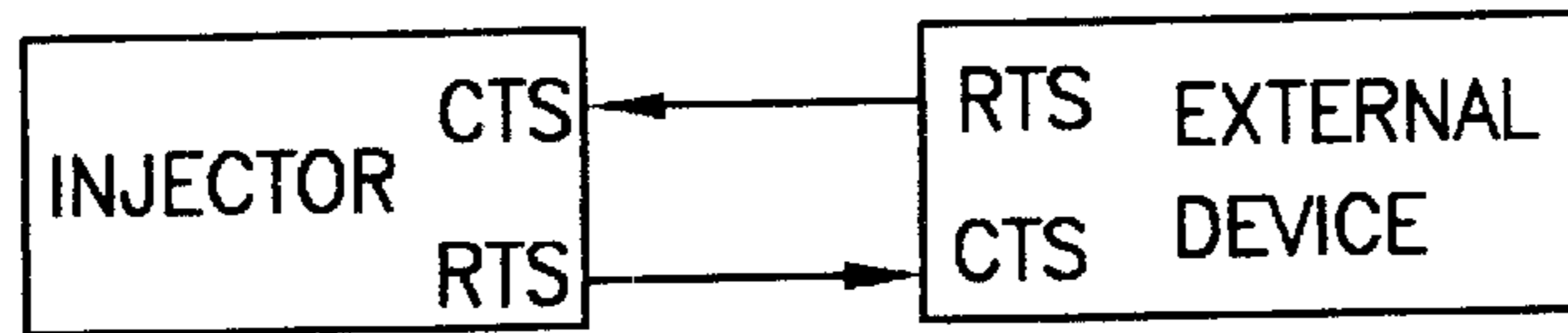


FIG. 5

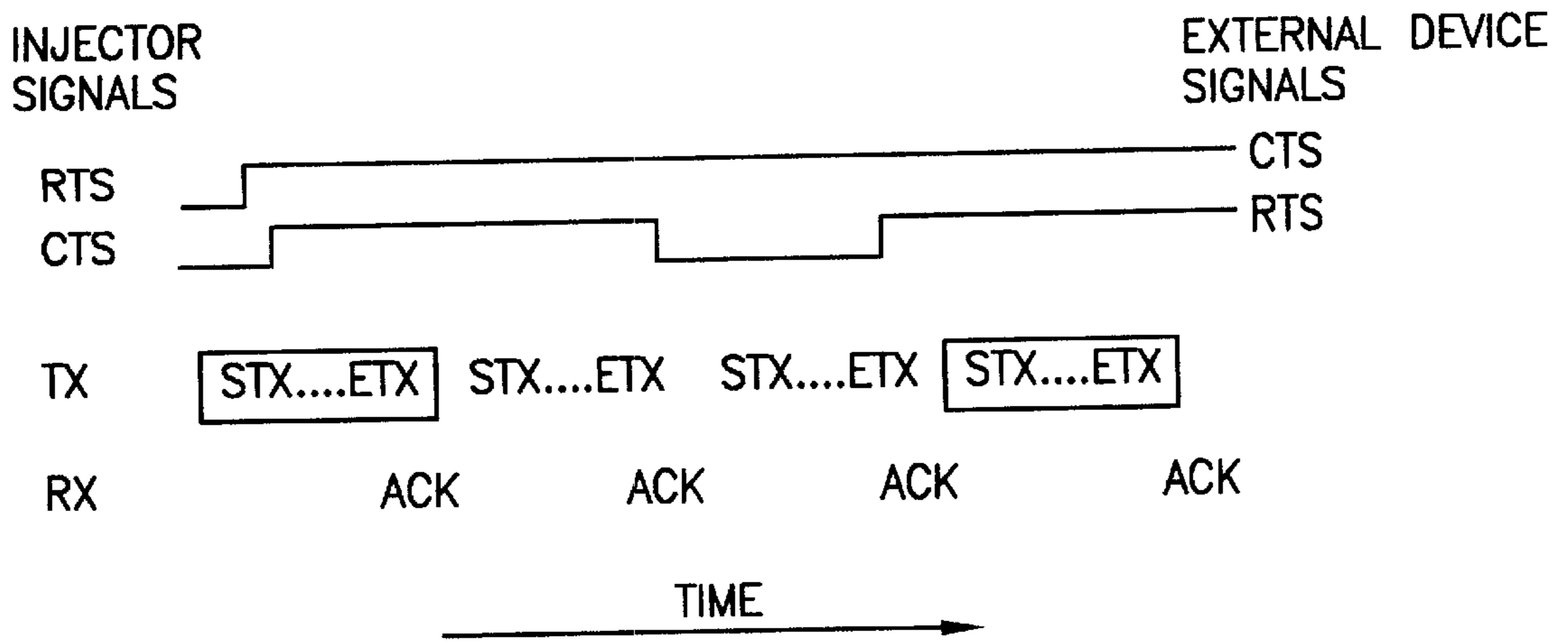


FIG. 6

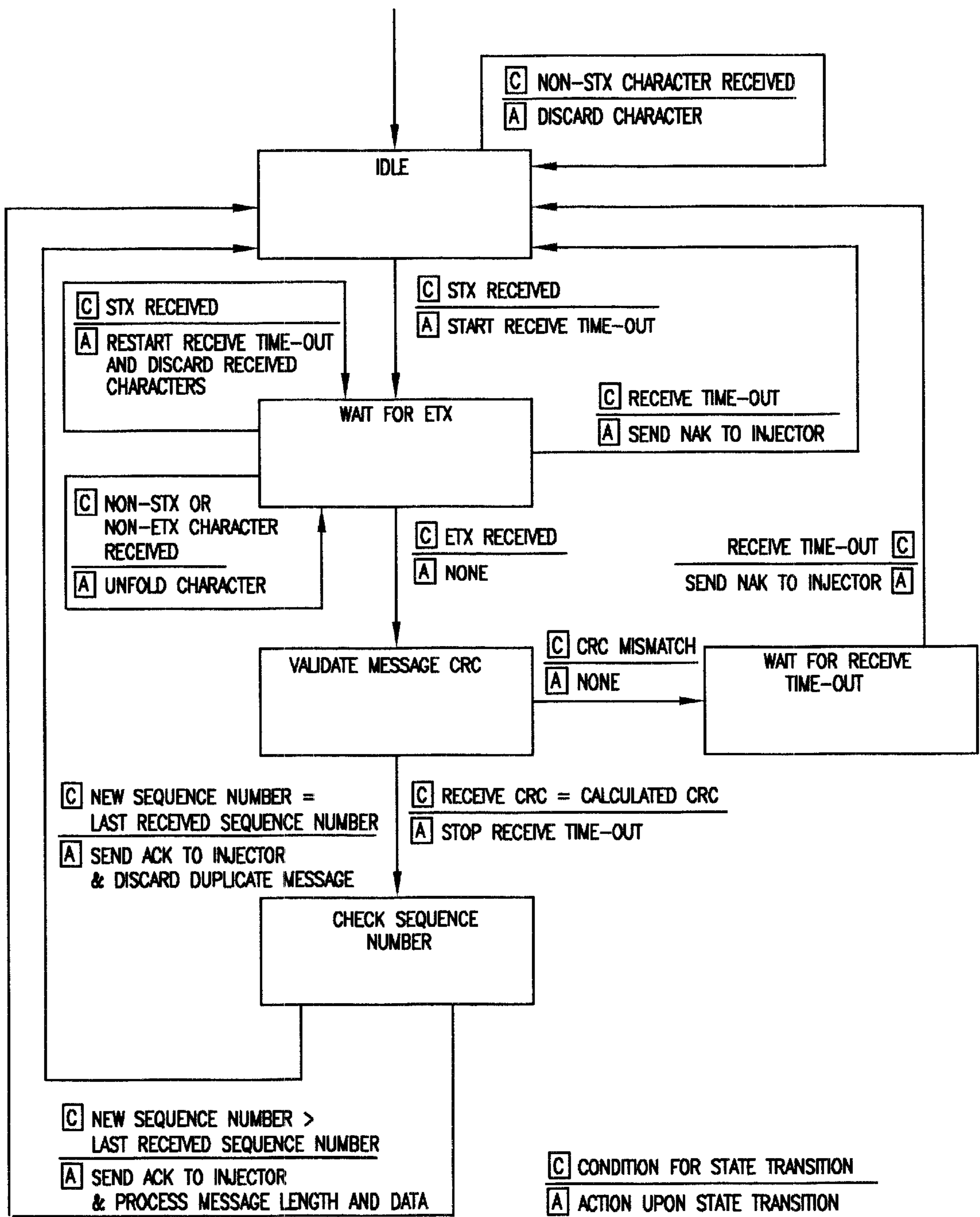


FIG. 7

DATA COMMUNICATION AND CONTROL FOR MEDICAL IMAGING SYSTEMS

RELATED REFERENCES

The present application is a continuation-in-part of U.S. patent application Ser. No. 09/197,773, filed on Nov. 23, 1998, which is a divisional of U.S. patent application Ser. No. 08/309,820, filed on Sep. 21, 1994, now U.S. Pat. No. 5,840,026, the disclosures of which are incorporated herein by reference.

FIELD OF THE INVENTION

The present invention relates generally to data communication and control, and, more particularly, to devices, systems and methods for communication of data and control based thereon in contrast-enhanced medical imaging systems.

BACKGROUND OF THE INVENTION

It is well recognized that the appropriate dose for many medications is related to a number of variables, including, for example, the size, weight, or physiologic state of the patient being treated. This variation is readily apparent from the different recommended doses many medications have for adults and children. The appropriate dose of contrast media for a given medical imaging procedure is equally dependent upon the size and weight of the patient being examined as well as additional factors.

Although differences in dosing requirements for medical imaging procedures have been recognized, many conventional medical imaging procedures, including angiographic, computed tomography, magnetic resonance and ultrasound imaging, continue to use pre-set doses or standard delivery protocols for injecting contrast media during medical imaging procedures. Although using fixed protocols for delivery simplifies the procedure, providing the same amount of contrast media to patients of widely varying size and weight can produce very different results in image contrast and quality.

Some of the shortcomings of existing procedures have been addressed and resolved as described in U.S. Pat. No. 5,806,519, issued Sep. 15, 1998, the disclosure of which is incorporated herein by reference. U.S. Pat. No. 5,806,519 discloses a contrast media delivery system that provides a source of contrast media sufficiently isolated from a patient undergoing an imaging procedure that the source of contrast media may be used on additional patients without concern for contamination. Additionally, the system is capable of adjusting contrast media concentration and other injection parameters during an injection procedure. The term "concentration" refers generally to the concentration of the image enhancing agents, particles or chemicals, in the contrast media.

The system of U.S. Pat. No. 5,806,519 incorporates a source of contrast media and, if desired, an admixture to dilute the media. If there is an admixture, then the contrast media preferably has a concentration which is the highest that would be used in an injection procedure so that the operator may combine the contrast media with the admixture and select virtually any concentration of contrast media desired for any given procedure. The concentration of the contrast media injected into a patient may be varied during the injection procedure by varying the ratio of admixture to contrast media. Each patient thus receives only the amount of contrast media necessary to provide a proper diagnostic image.

It has been recognized that the system of U.S. Pat. No. 5,806,519 is much more versatile and useful if the operator is able to select and adjust contrast media concentration and other injection parameters based on patient information and/or feedback received during the injection imaging procedures. U.S. patent application Ser. No. 09/197,773 and U.S. Pat. No. 5,840,026 disclose such systems. Those systems are capable of automatically choosing the appropriate concentration and injection rate for a given patient and are capable of automatically adjusting concentration and other injection parameters during an injection procedure based on feedback related to the resultant image quality.

Although significant strides have thus been achieved in the control of contrast-enhanced, medical imaging procedures, it remains desirable to develop improved devices, systems and methods for communication of data between devices and control based thereon in contrast-enhanced medical imaging systems.

SUMMARY OF THE INVENTION

The present invention provides several systems, devices and methods for producing a contrast-enhanced medical image of a patient. According to one embodiment, such a system includes generally a source of a contrast or enhancement medium and a pressurizing unit in connection with the source of contrast or enhancement medium to pressurize the contrast or enhancement medium for injection into the patient. The contrast medium or enhancement medium is also called the fluid medium, being a liquid, gas, or solid suspended in a liquid or a gas. The system also includes an energy source adapted to apply energy to a region of the patient and an imaging unit providing (preferably in real-time) an indication (for example, a visual or audible indication) of an internal state, condition or view of the patient based upon a signal resulting from the energy applied to the region of the patient. This signal is affected by a condition of the contrast or enhancement medium in the patient. The system also includes a control unit adapted to adjust the condition of the contrast or enhancement medium in the patient based upon the signal resulting from the energy applied to the region of the patient. The system may include general components or pieces of equipment manufactured by more than one company, for instance, an injector and an imager.

The condition of the fluid medium flowing into the patient can, for example, correspond to at least one parameter including, but not limited to, contrast medium concentration, flow rate of the contrast medium, timing of an injection, sequencing of more than one injection (of the same or different contrast media), injected volume of the contrast medium, injection pressure of the contrast medium or temperature of the contrast medium.

Although U.S. Pat. No. 5,840,026 describes generally the use of the pixel intensity of one or more portions or regions of the visual display of an imaging unit to control one or more conditions of the fluid medium within the patient, the present inventors have also discovered that direct use or modification of the raw signal used to produce the image may have certain significant advantages in controlling the imaging procedure. For example, a portion of the information available from the raw signal may be lost or distorted in various compression algorithms that are used to address the limited dynamic range of the displays that are available to show an image. The energy applied to the region of the patient may, for example, be sonic energy, in which case the raw signal is an acoustic intensity signal. This particular

method is sometimes called acoustic intensitometry or acoustic densitometry. The energy may, for example, also be penetrating radiation such as X-rays, or non-ionizing electromagnetic radiation such as light.

The present invention also provides another embodiment of a system for producing a contrast-enhanced medical image of a patient including a source of a fluid medium and a pressurizing unit as described above. As used herein, the terms "image" or "view" refer generally to an indication of an internal condition or state of a patient and are not limited to a visual display. The system also includes an energy source and an imaging unit to provide (preferably in real-time) an indication (for example, a visual display) of an internal view of the patient based upon a signal resulting from the energy applied to the region of the patient. In this embodiment, the system further includes a control unit adapted to adjust the condition (as described above) of the fluid medium in the patient to maintain at least one portion of the indication (for example, a portion of a visual display) at a desired level of intensity or enhancement. For example, the control unit can adjust the condition of the fluid medium flowing into the patient based upon the signal resulting from the imaging energy applied to the region of the patient. The system can also adjust the condition of the fluid medium flowing into the patient based upon measured intensity or density of the portion of a visual display. Such control of the imaging procedure assists in the study of, for example, lesions in the region of interest.

In another embodiment, the control unit adjusts parameters of the imaging unit to obtain an optimum or sufficient image based upon the concentration of the contrast enhancing agent flowing into the patient. Possible adjusted parameters include but are not limited to: the power in the signal sent into the patient, the time during which the energy is applied to the patient, the gain of the amplifier which receives the signal from the patient, or the speed at which the energy is scanned across the patient.

The present invention also provides a system for delivering an active substance (for example, a biologically active therapeutic substance or a diagnostic substance) to a patient including a source of a fluid medium that incorporates the active substance and a contrast agent. A pressurizing unit is in fluid connection with the source of fluid medium to pressurize the fluid medium for injection into the patient. The system also includes an imaging energy source adapted to apply imaging energy to a region of the patient and an imaging unit providing a visual display (preferably in real-time) of an internal view of the patient based upon a signal resulting from the imaging energy applied to the region of the patient. The system further includes a control unit adapted to control delivery of the active substance by adjusting the condition of the contrast agent in the patient.

In this embodiment, the active substance is preferably activated by activation energy from a source of activation energy. The activation energy and the imaging energy can be the same or different types of energy. In general, the concentration of the active substance in a region of interest will be directly proportional to the concentration or amount of contrast agent in the region of interest. The strength and duration of the applied activation energy can, for example, be adjusted based upon the signal resulting from the imaging energy or upon the resulting visual or other (for instance sound pitch, sound volume, numeric readout, or meter readout) indication provided to the user (preferably in real-time). The contrast medium and the active substance may be combined. For example, a therapeutic drug or a gene therapy may be contained in partially gas filled microspheres that are

ruptured by ultrasound energy beamed into a specific part of the body to activate the therapeutic drug or gene therapy.

The present invention further provides another system for producing a contrast-enhanced medical image of a patient similar to the systems described above. In this embodiment, however, the control unit is adapted to time injection of at least one discrete flow interval of fluid medium based, for example, upon at least one of a visual display or a signal resulting from application of the imaging energy to a region of the patient (both may be used simultaneously). In one embodiment, the discrete flow interval may be a bolus of fluid medium.

Preferably, unidirectional or bi-directional communication and optionally control between devices of the present invention is enabled through use of a control/communication interface to which each of the devices of the imaging/injection system can be connected. This interface mating point can be located on a separate device or can be incorporated into one of the other devices (for example, into a controller for the pressurizing unit or injector). Suitable communication methods include data transmission, preferably digital, over wires, fiber optics, or via conducted or broadcast electromagnetic radiation (for example light and RF) or ultrasonic radiation.

The communication interface of the present invention is a substantial improvement over previous systems that merely communicated timing information via relay closures. In the present invention, data transmission includes information sent between devices regarding operating parameters, operator input, device status information, and/or control sequencing. In addition, data transferred from one device to another device in the present invention can be used to enable active control of the receiving device from that data. For example, during the injecting state, data transmission from an ultrasound imager can be sent to an injector to enable active control of the injection flow rate based on the data received. Currently available systems merely relay analog closures between an injector and an imager to communicate the timing of certain states and are not used for data transmission or communication. In such systems, the relay closure causes the injector or imager subsystem to begin executing a preset program. In the present invention on the other hand, data on the status of one subsystem is communicated to the other subsystem for use by that subsystem. The receiving subsystem may alter its operation based upon this information, even to the extent of being programmed by the transmitting subsystem. Moreover, digital communication enabled by the present invention allows much more information to be conveyed than a simple relay closure.

Preferably the communicating devices use the same protocol so that the information being communicated does not need to be converted. Note that it is also possible for the devices to support multiple communication protocols which may be selected by the user or selected by automatic negotiation between the communicating devices.

In addition, the present invention provides methods for adjusting the condition of a fluid medium (as described above) during an imaging procedure. In one embodiment, the method includes pressurizing the fluid medium for injection into the patient. Further, the method includes supplying energy to a region of the patient and indicating (preferably in real-time) an internal view (that is, a condition or state) of the patient based upon a signal resulting from the energy applied to the region of the patient. This signal is affected by a condition of the fluid medium in the patient as described above. The method further comprises the step of

adjusting the condition of the fluid medium in the patient based upon the signal resulting from the energy applied to the region of the patient.

In another method, the condition of the fluid in the patient is adjusted to maintain at least one portion of the indication (for example, a portion of a visual display) at a desired level of intensity/density or intensity/density profile over time. For example, the control unit can adjust the condition of the fluid medium in the patient based upon the signal resulting from the energy applied to the region of the patient to provide enhancement in the region of interest that is constant or varies within some acceptable range. The system can also adjust the condition of the fluid in the patient based upon measured intensity of the portion of a visual display or measured intensity of an acoustic indicator. As discussed above, such control of the imaging procedure can assist, for example, in the study of lesions in the region of interest.

The present invention also provides a method for delivering an active substance to a patient, comprising the step of controlling delivery or administration of the active substance by adjusting the condition of the contrast agent in the patient. As described above, the active substance is preferably activated by activation energy from a source of activation energy. The activation energy and the imaging energy can be the same or different types of energy.

The present invention also provides for structured communications in the situation where the control unit functions are shared between two or more pieces of equipment.

For example, the present invention provides a system for control of an imaging procedure comprising: a source of a contrast or enhancement medium; a pressurizing unit in fluid connection with the source of contrast or enhancement medium to pressurize the contrast or enhancement medium for injection of the contrast or enhancement medium into the patient; an imaging energy source adapted to apply imaging energy to a region of the patient; an imaging unit providing an indication (for example, a visual display) of an internal view of the patient based upon a signal resulting from the imaging energy applied to the region of the patient, the signal being affected by a condition of the contrast or enhancement medium in the patient; and a data communication interface between at least the pressurizing unit and the imaging unit to enable exchange of data between the pressurizing unit and the imaging unit. Preferably, the exchange of data is bi-directional. Moreover, the data is preferably in digital form.

The present invention also provides an injector system for producing a contrast-enhanced medical image of a patient in cooperation with an imaging system. As described above, the imaging system applies energy to a patient and produces an image or a measurement of a region of interest in the patient from a signal resulting from the applied energy. The injector system preferably comprises: a source of a contrast or enhancement medium; a pressurizing subsystem in connection with the source of contrast or enhancement medium to pressurize the contrast or enhancement medium for injection into the patient; an injector control unit for controlling said pressurizing subsystem; and a communication interface to exchange data between the injector system and the imaging system.

The injector system may communicate information from the injector system to the imaging system. Likewise, the imaging system may communicate data or information from the imaging system to the injector system. Such communication can be unidirectional or bi-directional. The injector control unit may modify one or more parameters of the

injection based upon the data or information from the imaging system. The imaging control unit may also modify one or more parameters of the imager unit based on the data or information sent from the injector. The injector system may further contain an electrically and/or physically isolated (wireless) interface through which the communication interface exchanges data or information with the imaging unit.

The present invention also provides an imaging unit for producing a contrast-enhanced medical image of a patient in cooperation with an injector system. As described above, the injector system pressurizes a contrast or enhancement medium for injection into the patient. The imaging unit preferably comprises: a source of energy to be applied to a region of interest in the patient; a display to provide an image based upon a signal resulting from the imaging energy applied to the region of the patient; an imaging control unit for controlling the imaging unit; and a communication interface to exchange data or information between the injector system and the imaging unit.

Communication of data/control may be from the injector system to the imaging unit or visa versa and either uni- or bi-directional. The imaging unit preferably further contains an electrically and/or physically isolated interface through which the communication interface shares information with the injector system.

The interface is preferably in communicative connection with at least the pressurizing unit and the imaging unit to enable sharing and exchanging of data between the pressurizing unit and the imaging unit.

The above and other systems, devices and methods of the present invention, and their attendant advantages, will become even more apparent to one skilled in the art upon consideration of the following detailed description in combination with the accompanying drawings.

BRIEF DESCRIPTION OF THE DRAWINGS

FIG. 1 illustrates an embodiment of an injection and imaging system of the present invention using ultrasonic energy to display an image of an internal region of the patient.

FIG. 2 illustrates an embodiment of a data communication unit of the present invention.

FIG. 3 illustrates another embodiment of an injection and imaging system of the present invention.

FIG. 4 illustrates an example of internal transmit and receive sequence numbering.

FIG. 5 illustrates an embodiment of a connection between an injection system and external device, showing physical layer control signals.

FIG. 6 illustrates an embodiment of the communications protocol between an injector system and external device, showing message flow control.

FIG. 7 illustrates an example of a state transition diagram of the receive processing to be performed by an external device communicating with an injector subsystem of the present invention.

DETAILED DESCRIPTION OF THE INVENTION

The present invention will be discussed, in part, with reference to a model injection and imaging system 10 illustrated in FIG. 1, in which ultrasonic energy is applied to a patient 20 to display an image of an internal region of patient 20. However, it should be understood that the present

invention is also applicable to other imaging modalities, including x-ray, computed tomography, magnetic resonance imaging, nuclear imaging, and other modalities in which energy is released within or transmitted through a portion of the human body, animal, or other object to be indicated or imaged.

Ultrasound imaging creates images of the inside of the human body by broadcasting ultrasonic energy into the body and analyzing the reflected ultrasound energy. Differences in reflected energy appear as differences in gray scale or color on the output images. As with other medical imaging procedures, contrast enhancing fluids (often referred to as contrast media) can be injected into the body to increase the difference in the reflected energy and thereby increase the gray scale or color contrast displayed in the image (that is, the image contrast) viewed by the operator.

For ultrasonic imaging, the most common contrast media contain many small bubbles (sometimes referred to as microbubbles). The difference in density of bubbles when compared to water, and thus their difference in sound transmission, makes small gas bubbles excellent means for scattering ultrasound energy. Small solid particles can also serve to scatter ultrasonic energy. Such particles are typically on the order of 1 to 10 microns (that is, 10^{-6} to 10^{-5} meters) in diameter. These small particles can pass safely through the vascular bed, and in some cases, traverse the pulmonary circulation. Contrast agents are also used for non-vascular applications such as assessment of tubal patency in gynecological applications.

Contrast media suitable for use in ultrasound are supplied in a number of forms. Some of these contrast media are powders to which liquid is added just before use. The powder particles cause gas bubbles to coalesce around them. The powder must be mixed with a liquid, and the mixture must be agitated with just the right amount of vigor to get the optimum creation of bubbles. Another type of contrast medium is supplied in a liquid form that requires hypobaric or pressure activation. A third type of contrast medium is a liquid that is agitated vigorously. There are no solid particles to act as nuclei, but the liquid is a mixture of several liquid components that make relatively stable small bubbles. A fourth type of contrast medium uses "hard" spheres filled with a gas. These contrast media are typically supplied as a powder that is mixed with a liquid. The goal is to suspend the spheres in the liquid without breaking them. Even though such spheres have a shell that is hard compared to a liquid, they are very small and relatively fragile. It is also possible for the solid particles themselves to act to scatter ultrasonic energy, but the acoustical properties of the solid spheres are not as different from water as those of a gas. Entities such as microbubbles, microspheres and solid particles suitable to enhance ultrasonic imaging contrast are referred to herein as contrast enhancement agents. For X-ray based imaging, atoms, molecules or solid particles generally absorb X-rays at a higher rate than blood and gas or gas-filled structures generally absorb X-rays at a lower rate than blood.

Contrast enhancement agents also enhance other modes of ultrasonic imaging. For example, when the microbubbles, microspheres or particles are carried along in the blood stream, the reflected energy is Doppler shifted. This Doppler shift allows an estimation of the speed of blood flow. Bubbles can also be excited so that they radiate ultrasonic energy at an harmonic of the incident ultrasonic energy. This harmonic imaging is only possible with the use of contrast medium.

After mixing/preparation as described above, the contrast medium is drawn into a syringe or other container for

injection into the patient. Typically, the fluid is injected into a peripheral vein in the arm of the patient, although it may be injected into other body cavities for specific imaging procedures, for instance the intestinal tract or the female reproductive tract. When injected into the bloodstream, the blood dilutes and carries the contrast medium throughout the body, including to the area of the body being imaged (that is, the region-of-interest or ROI).

It is becoming more common for a microprocessor-controlled, powered injector to be used for injecting the contrast medium. The use of such powered injectors provides the benefit of maintaining a consistent flow over an extended time period, thereby providing a consistent amount of contrast medium in the blood stream. For ultrasound, if there are too few contrast enhancement agent particles per unit volume in the flow, however, an insufficient image contrast may result and thereby prevent an accurate diagnosis from being made. If too many contrast enhancement agent particles are present, on the other hand, excess energy is reflected, thereby resulting in blooming or saturation of the ultrasound image.

Thus, although a power injector can inject contrast medium at a constant flow rate (volume per unit time), there typically must be a generally constant number or amount of contrast enhancement agent particles per volume of fluid injected to provide a constant image contrast. If the concentration of particles changes, the image contrast will be degraded. There are many reasons why the number of contrast enhancement agent particles per volume of a certain contrast medium (and thereby the image contrast) can vary during an injection procedure. For example, density differences between the contrast enhancement agents and the fluid medium may result in separation. Moreover, the initial mixing may not have resulted in a homogeneous dispersion or suspension. Likewise, bubbles or microspheres of certain contrast media can be destroyed under conditions experienced in mixing, storage or delivery of the contrast media. Additionally, a number of conditions or parameters of the injection fluid other than the concentration of the contrast enhancing agents therein can be changed. For example, the injection flow rate, the volume of medium injected and the injection pressure can be changed in a continuous or discontinuous manner. In addition most newer ultrasound enhancement agents and most existing X-ray enhancement agents pass through the pulmonary circulation and can circulate more than once through the body, so that they build up in the patient's blood pool. The extent to which this happens differs from patient to patient. This is another reason for having feedback from the imaging system to the injector to control the injection.

Referring to FIG. 1, injection and imaging system **10** delivers a fluid medium including a suspension of microbubbles **30** suspended in a fluid carrier to patient **20**. Certain components of system **10** are discussed in detail in U.S. application Ser. No. 09/267,238 entitled APPARATUS AND METHOD FOR CONTROLLING CONTRAST ENHANCED IMAGING PROCEDURES, filed on Mar. 12, 1999, the disclosure of which is incorporated herein by reference.

In the embodiment of FIG. 1, a syringe or pressurizing vessel **40** drives fluid through a fluid path element **50**. Fluid path element **50** may pass through a bubble or contrast enhancement agent concentration regulator **60** that can affect the enhancement properties of the agent by, for example, selectively destroying microbubbles **30**. Bubbles **30** may be destroyed, for example, by insonating the fluid with ultrasound energy, generating local temperature or pressure changes in the media or by some types of mechanical agitation.

The ability to destroy microbubbles **30** in the fluid path may enable better control of the imaging procedure. Because mechanical resonance of a bubble wall is a function of bubble size, for example, it may be possible with the proper power and frequency settings to selectively decrease the concentration of bubbles of a certain size. Such selective destruction allows control of the bubble distribution. Microbubbles **30** can also be destroyed in the fluid path as part of a strategy to control contrast agent concentration. It is also possible to reduce the amount of contrast enhancing agent flowing into the patient simply by reducing the flow rate of the media.

Fluid path **50** continues through a concentration sensor **70** suitable to measure the concentration of microbubble **30** in the fluid medium. Such sensors are discussed in detail in U.S. application Ser. No. 09/267,238 referred to above.

Fluid path **50** then continues on through a patient interface **80** to an injection site on patient **20**. The output of sensor **70** preferably passes to a control/communication interface **90** which may, for example, be in communicative connection with or combined with a processor unit **92** suitable to, among other things, process signals corresponding to, for example, concentration data from sensor **70** and to send control signals based upon such incoming data to other components of system **10**. A control signal corresponding to the concentration data may be sent from processing unit **92** via interface **90** to any number of devices in delivery system **10** including, for example, an imaging unit such as an ultrasound scanner **100**. The control signal can be used, for example, to adjust the image (for example, by increasing or decreasing the power in the signal sent into the patient, the time during which the energy is applied to the patient, the gain or other machine settings of the amplifier which receives the signal from the patient, or the speed at which the energy is scanned across the patient), to provide concentration information as part of documentation and/or to assist with diagnosis during the imaging procedure. Information can also preferably be sent from imaging unit **100** to processing unit **92** to, for example, control other devices such as, for example, an agitation/preparation mechanism **110** (including, for example, a mechanical stirrer **114**), a powered injection control unit **120** and/or a concentration regulator **60**.

Agitation/preparation mechanisms suitable for use in the present invention are disclosed in U.S. application Ser. No. 09/267,237 entitled AGITATION DEVICES AND DISPENSING SYSTEMS INCORPORATING SUCH AGITATION DEVICES, filed on Mar. 12, 1999, the disclosure of which is incorporated herein by reference. In that regard, several commercial agents now available require some kind of agitation or mixing for preparation. These include, but are not limited to: ALBUNEX and OPTISON available from Molecular Biosystems, Inc. of San Diego, Calif.; and LEVOVIST available from Schering AG of Berlin, Germany.

The data communication and control devices and systems of the present invention can be used during initial preparation of the medium during an injection procedure to effect a desired result, or after an injection to review performance and status. In that regard, certain contrast media, such as LEVOVIST, are known to precipitate or separate over time in some of the higher concentrations (for example, 400 mg/ml). Concentration measurement can be used to detect such separation and restart or increase agitation to help reduce precipitation effects.

Concentration sensor **70** can be located anywhere on the fluid delivery path from syringe or pressurizing vessel **40** to

patient interface **80**. If concentration sensor **70** is located near the patient injection site at the end of fluid path **50**, one can better account for microbubble degradation effects from shear rate, temperature and other delivery effects. Since sensor **70** preferably does not require direct contact with the contrast enhancement agent, a coupling piece can be made as a disposable part of a tubing set for convenience in attaching sensor **70** and or concentration regulator **60** and to maintain sterility with patient interface **80**. The sensing region is preferably a known volume or area of the contrast media surface located on fluid path **50**.

Other sensors or sensing equipment can be in communicative connection with interface **90**. For example, a sensor or sensors (not shown) can be part of system **10** to provide direct measurements of the concentration of the fluid medium and/or the contrast enhancement agents within the patient's body. Another example is bar code readers which input information about the contrast media including but not limited to volume, concentration, manufacture date, manufacturer. Moreover, interface **90** can be in unidirectional or bi-directional communicative connection with a sensing and/or measuring device **130** that provides data on one or more physiological conditions of the patient. For example, device **130** can be a heart rate or blood pressure monitor.

Previous injector systems have contained an electrocardiograph (ECG) that is used to set start of injection trigger points synchronized with various portions of the cardiac cycle. In the present invention, other equipment such as heart rate or blood pressure monitors may be used to provide patient physiological data to control the injection or imaging process, especially during an injection. For example, during ultrasound cardiac stress echo imaging, vasodilator and vasoconstrictor drugs are administered to increase and decrease cardiac output load. Connected sensors for monitoring heart rate and blood pressure may be used to initiate, terminate or adjust the injection process when these parameters reach certain levels. The output from these sensors can also be used to adjust other aspects of the imaging process, such as scanner settings. Unlike prior systems, the present invention allows isolation, digital communication, bi-directional or two-way communication and/or communication with an imaging system.

In addition, it may be useful to control the delivery of some therapeutic or diagnostic agents during contrast imaging based on display or signal information from the imager as a result of the imaging procedure. For example, during ultrasound cardiac stress echo procedures, a drug, Dobutamine, which is a cardiogenic or cardiovascular stressor agent, is often used to increase cardiac stress so that cardiac parameters can be measured and quantified. (Dobutamine is a synthetic derivative of dopamine, characterized by prominent inotropic but weak chronotropic and arrhythmogenic properties.) An injection system may be used to control the delivery of the stressor drug during an enhanced procedure based on visual information from the imager, such as peak flow rate in a vessel from a Doppler image, the intensity of a perfused tissue region, vessel geometry measurements, or heart chamber geometry measurements during various portions of the cardiac cycle. The delivery of such a drug can also be controlled by the signal at the imager before it is generated into a display, for example, using acoustic intensitometry. This concept can be extended to other imaging modalities and to the administration of other therapeutic or diagnostic drugs or substances.

Interface **90** can also be in communicative connection with an information storage and processing unit **140** of a

hospital or other care-giving organization to, for example, receive information regarding a patient to assist in control of injector controller **120** or to transmit information regarding an imaging procedure from any other device of imaging system **10** for storage and/or processing by hospital processing unit **140**.

Interface **90** can also be in communicative connection with an operator interface **150** so that the operator(s) of imaging system **10** may receive information about the patient and the components of imaging system. Data, such as control parameters or algorithms, can also preferably be sent from user interface **150** to interface **90**. It may be preferable to physically partition the operator interface so that some simple injection related information such as volume remaining is available near the syringe. It is preferable that all injection and scanning information be available so that the operator can quickly access relevant information during the imaging procedure. Currently, ultrasonic imagers use a display that combines the image and informative text around the image. Two different displays may be used for cost reasons or for ease of use. Information that may be displayed for operator consideration includes but is not limited to: time since the fluid was initially prepared, information such as brand or initial concentration read from contrast media packaging, initial injection parameters for a single or multilevel injection, and the status of the injector controller (for instance, disarmed, armed, injecting).

Operator interface **150** also preferably incorporates operator input devices including but not limited to keys, a track ball, a mouse, a joystick, a voice recognition system, and/or a touch panel as known in the art. These input devices allow the operator to make initial settings or to input information about the patient or the imaging procedure which the processor **92**, the controller **120**, or the imager **100** can use to prepare for or conduct the imaging procedure. The operator interface also enables the user to ready the injector for injection (arm the injector) or to disarm. It can also allow the operator to confirm that the proper procedure has been followed in eliminating gross air from the fluid path elements. The operator interface allows the operator to determine information used from or sent to other components such as but not limited to optional use of patient information such as patient physiologic parameters through external device interface **130** or output of selected information to the hospital processor **140**.

An embodiment of interface **90** will be discussed in the context of communication with injector control unit **120**. FIG. 2, for example, illustrates an embodiment of interface **90** in communicative connection with injector control unit **120** via cabling **94**. Interface **90** can preferably be connected to other devices of system **10** via any wired or wireless media for carrying the information (for example, RS-232, parallel, RS-485, LAN, etc.) using any unidirectional, paired unidirectional, or bi-directional networking or communication protocol (for example, TCP/IP, IPX/SPX, etc.) as preferably selected by an operator. Refer to Table 3. for a summary of some of the possible communication connections and protocols.

Data communication and control can be effected with use of interface **90** in part, for example, through definition of various operational states of the components of system **10**. For example, a ready/armed state can be defined as the state of injector controller **120** prior to performing an injection. This state may be set upon completion of specific diagnostic functions and indicates that injector controller **120** is ready and awaiting a signal to begin an injection. An idle/disarmed state can define an idle state of injector controller **120** during

which programming operations are performed. Preferably, the idle/disarmed state is the power-up default state and the state that injector controller **120** returns to after performing any injection sequence. There can be an armed multi-inject state which allows a series of injections before returning to the idle state. Injector controller **120** is preferably always in the idle/disarmed state when connecting to or disconnecting from patient **20**. A hold state of injector controller **120** is preferably entered after the operator or another system component has temporarily suspended the injection. Injector controller **120** may, for example, remain in this state for a period of time before disarming while awaiting a signal to resume the injection. An injecting state defines the state of injector controller **120** while performing an injection. This state is entered typically only from a ready/armed state.

A set of injection conditions or parameters are preferably entered into the system (for example, by an operator via operator interface **150**) to define certain criteria of how an injection is to be performed. These conditions or parameters can include, for example, volume, flow rate, number of phases, fluid source or type (where A and B are two types of fluid), and whether there is a scan delay or injection delay associated with the programmed injection (that is, delivery of fluid according to a programmed injection profile).

Operational and state parameters are preferably provided between components of system **10** via interface **90** (for example, between injector controller **120** and imaging unit **100**) at an update rate sufficient to allow effective real-time control of the imaging procedure (for example, to allow imaging unit **100** to produce a clear image of the injection process). For peripheral venous injections control does not need to be instantaneous because there is a variable delay from the time the fluid is injected into the patient to the time it reaches the imaging site, as well as a delay from when the fluid passes the concentration sensor **70** and the time it flows into the patient. A second function of the interface **90** is to allow control parameters to be passed between components of imaging system **10**.

Typical operating parameters include, for example, programmed infusion rate, programmed bolus rate, programmed bolus volume, flow rate, total injected volume, volume remaining, bolus ready, infusion ready, bolus running, infusion running, low volume, infusion pending, bolus complete, flow profile rise/fall rates, injection aborted/canceled, error codes as well as internal diagnostic information and injector machine state. Typical control parameters may include, for example: hold injection, start injection, start scan, stop injection, stop scan, gated operations (Start/Stop) in relation to some patient parameter such as breathing or ECG, reset injector, programming commands, preparation commands, loading commands, or any other injector functions that could be activated normally by the user or by other means.

The following is an example of a data communication protocol for use with interface **90**. Imaging unit **100** may, for example, include an ultrasonic scanner that operates in conjunction with an injection system (including injector controller **120** and pressurizing syringe **40**) to inject a contrast agent to enhance a targeted area of concern or region of interest of patient **20**. Injector controller **120** and imaging unit **100** may be interconnected via interface **90** (for example, including an RS-232 serial interface) using standard control lines Request to Send (RTS) and Clear to Send (CTS) to manage the data flow. A simple protocol may be defined wherein injector controller **120** provides all of its operating parameters via interface **90** periodically, for example every 500 milliseconds, as long as the CTS line

from imaging unit **100** is set (injector controller **120** signals its readiness to imaging unit **100** by setting RTS). After receipt of a message imaging unit **100** may simply acknowledge data packets received from injector controller **120**. Imaging unit **100** can also send control commands or in a separate message as necessary to control the injection/ imaging process. Injector controller **120** may simply acknowledge command packets received from imaging unit **100** and may perform the requested action if it is safe to so.

In one sequence of events, an operator sets up injector controller **120** by following all required procedures to the point where injector controller **120** is in an 'Armed' or 'Ready' state. At this point the operator can concentrate fully on operation of imaging unit **100**. When ready, the operator can use a command from the imaging unit **100** to injector controller **120** (via interface **90**) to begin a programmed injection. During the programmed injection and scan, the injector controller **120** is preferably providing a rapid update of its process and status. At any time, the operator preferably can abort the injection (from either the injector or imaging system or take over active control, for example, adjust the flow rate.

Interface **90** thus facilitates an integrated control feature between devices of injection and imaging system **10**, permitting enhancement of diagnostic information. System **10** gives an operator the provisions to manage many or all of the components of system **10** (and, thereby, all facets of the injection and imaging procedure) from the most convenient place. Indeed, a single operator can easily perform a contrast-enhanced scan using the procedural ease afforded by the integrated approach of system **10**. Any number of quantification algorithms that expand the diagnostic capability of the system can utilize the information provided over interface **90** to maximize the quality and diagnostic utility of the images.

For example, there are numerous applications for an ultrasound injector with a bi-directional data and control communications interface **90**, especially in the areas of closed loop control of injector controller **120** based on an acquired signal or image. Such applications may be divided broadly into two classes, diagnostic and therapeutic.

As a diagnostic application, a communication interface **90** between injector controller **120** and scanner **100** can be used to detect lesions and tissue characteristics through differences in tissue vascular concentration or perfusion. The technique is useful as a method for tumor detection and identification. Closed loop control can, for example, be used to provide relatively constant enhancement of tissue within a region of interest to help identify tumors in those area.

Video density, video intensity, acoustic intensity or acoustic Doppler intensity information (that is, raw acoustical data) from scanner **100** can be used to control the injection profile (for example, as a way to control injected enhancement agent concentration) to provide the desired time response and level of enhancement. An algorithm in processor unit **92** or control unit **120** can accommodate for delays between a change in an injection parameter such as flow rate and the appearance of the change at the imaging site. Because tumors are often hyper-vascular or hypo-vascular relative to surrounding normal tissue, tumors can be identified by differences in enhancement level or enhancement level changes over time. A similar technique can be used to observe and quantify tissue perfusion during contrast enhanced myocardial echocardiography or other perfusion studies.

Closed loop control can be used to control the concentration of the enhancement agent in a region of tissue. The

level of enhancement required, the level of enhancement or the time response of enhancement increase or decay can be used to observe tissue perfusion, since the enhancement agent will be present in the small vessels that perfuse the region of interest. This technique can also be combined with the use of vasoconstrictor/vasodilator drugs, physical stress, or other means to induce cardiovascular stress to make changes in perfusion more pronounced or detectable.

Another diagnostic application for interface **90** is to control parameters such as concentration, flow rate or time or timing of injection based on data of a physiologic parameter, such as heart rate, provided by external device **130**. An example of a procedure is closed loop drug delivery of cardiac stress agents while imaging. In this technique, a stressor agent is injected and controlled based on heart rate. Once a target heart rate is achieved it may be useful to synchronize and control the level of injected enhancement agent, or to provide bolus injection control for diagnostic purposes. In that regard, information from imaging unit **100** can be used to control the delivery of the enhancement agent. Interface **90** can also be used to terminate an injection if a certain level of enhancement has not been reached within a certain time period, preventing waste of the imaging agent and saving procedure time. Or it can simply warn the operator that conditions are outside some optimal window and let the operator take appropriate actions. This technique can be generalized to injection communication and control based on information from other external equipment or patient physiological parameters.

Information from the concentration sensor **70** can be used to affect the injection parameters, or it can simply be communicated via interface **90** to the operator interface **150** so that the operator has access to the concentration information during the imaging procedure. The operator is the one who takes action to either adjust the injection, stop the procedure or take other appropriate action. This has a potential advantage in achieving quicker regulatory approval for the manufacture and sale of the device. This allows the operator to make the decision vs. the system taking action automatically which may require more validation effort. Also data from concentration sensor **70** could be transmitted to the imager **100** directly or after processing in the processor **92** so that parameters of the imaging unit mentioned elsewhere can be adjusted to achieve a satisfactory diagnostic image in spite of changes in concentration of the contrast enhancing agent.

There are also many therapeutic and other applications of imaging system **10**. For example, enhancement agents may be used as a means for site specific imaging and for activated drug or gene therapy delivery. For example, ImaRx of Tucson, Ariz. has an agent under development, MRX-408, based on their previous agent DMP-115 (DEFINITY), a multilipid encapsulated perfluoropropane and air echo enhancing agent that incorporates site specific binding for the detection of thrombus. Some ultrasound and x-ray imaging agents are being developed that release drugs or are "activated" when insonated or bombarded with x-ray energy. Image video intensity, acoustic intensity, acoustic Doppler intensity or other image information can be used with these agents to track and control the concentration of an agent within a region over time. Coupled with injector controller **120** and interface **90** for injection control, the delivered agent concentration or total agent dose can be controlled for maximum therapeutic effect and to prevent overdose.

For example, a certain drug delivery imaging agent can be injected under control of imaging system **10** so that the observed enhancement is stabilized at a target level for a

certain period of time. During this time, the drug of the agent can be activated within the region of interest using, for example, ultrasound or x-ray energy. Once the time period of sufficient enhancement expires, the injection can be terminated by control from imaging unit **100**. Information on the injection delivery history (e.g., the flow rate profile) and total dose (e.g., total volume delivered) can be sent to the imaging unit **100** or hospital processing unit **140** for record keeping and documentation purposes. Other methods, such as using the time integral of image pixel intensity, can be used to measure and control the total delivered dose of the agent. With microbubble-based delivery of active substances in ultrasound, information from the imaging unit **100** can also be used to make sure that all of the agent within a certain region has been completely activated before ending the injection, maximizing the delivered dose within the region of interest. Other agents may become available in the future that are activated by electromagnetic (especially optical), magnetic, or nuclear radiation. All such activation energy sources are suitable for use in the present invention.

Another application of real-time data and control interface **90** is for control of pulsed flow for therapeutic applications. In pulsed flow thrombolysis, for example, high flow rate but short duration bolus injections are used to both mechanically disrupt or ablate and chemically dissolve thrombi in vessels. Image information regarding the progress of clot dissolution can be used to communicate changes to the injection system flow rate, agent concentration, pulse duration and other fluid delivery parameters. While not shown in the figure, a duplicate injector with duplicates to syringe **40** and control unit **120** could be injecting the therapeutic agent into the patient and cooperating through interface **90** with simultaneous or alternating injections of therapeutic agent, flushing agent (such as saline) and imaging agent. In some applications it can be advantageous to control the injection of three (3) or more fluids by using multiple injection mechanisms in communications with each other through interface **90**.

In certain therapeutic injections, it is desirable to synchronize the delivery of short boli of therapeutic fluid with that portion of the cardiac cycle that ensures turbulent mixing with blood. Without turbulent mixing, these toxic fluids can be entrained in streamlines of laminar flow which carry the fluid beyond the area of intended application. In this case, the fluid is typically introduced by a peripheral arterial access. A precise volume of fluid must be injected by breaking it up into small volume packets timed to the cardiac cycle. A communications link to injector controller **120** can be used to control the timing of fluid delivery in this application.

FIG. 3 illustrates an example of a block diagram that partitions the functions described above between separate injection subsystems and imaging subsystems. This embodiment may, for example, accommodate a situation in which the various subsystems are separately manufactured by the same or different manufacturers and/or a situation in which the subsystems require separate regulatory approvals.

In FIG. 3, the unpartitioned components retain the same numbers as in FIG. 1. For those that are partitioned between the two subsystems, those partitioned into injector subsystem **15a** of the system generally include a suffix "a" and those partitioned into imager subsystem **15b** generally include a suffix "b." Component **90c**, is the interface connection between the two subsystems.

Injector subsystem **15a** includes generally those elements for the control of the fluid flow. Injector subsystem **15a**

includes, for example, concentration sensor **70** and external device **130**. Injector processor unit **92a** takes and receives information from an injector operator interface **150a** and the other components through injector interface **90a**.

Imager subsystem **15b** includes generally elements for the control of the imaging energy and the creation of an image or measurement as discussed above. Imager subsystem **15b** includes, for example, a processor unit **92b** that takes and receives information from an imager operator interface **150b** and the other components through imager interface **90b**.

It is possible for the subsystem components to be partitioned in other ways. For instance, concentration sensor **70** might be part of the imager subsystem **15b**. The same applies to external device **130**. FIG. 3 shows both subsystems having separate communications to the hospital information system **140**. It may be that only one of the subsystems communicates with the hospital interface **140**. Other partitioning is possible as well.

All communications between the two subsystems preferably take place through subsystem interface **90c**. A benefit of this partitioning is the isolation of the need for the most stringent error control and verification to the operation of subsystem interface **90c**. Communications within each subsystem can be more easily tested as a self-contained unit.

Present imaging systems use a simple relay closure as an interface between subsystems so that one subsystem can tell the other to start a preprogrammed function. By having a subsystem interface **90c** transmit information, many of the above listed benefits are possible.

In addition to timing information communicated via relay closures by prior systems, the data transmission enabled by the present invention includes, for example, information sent between devices regarding operating parameters, operator input, device status information, and control sequencing. In addition, data transferred from one device to another device can be used to enable active control of the receiving device from that data. For example, during the injecting state, data transmission from an ultrasound scanner can be sent to an injector to enable active control of the injection flow rate based on the data received.

The discussion below provides a list of example capabilities possible with interface **90c** between the subsystems of the present invention. Many of these have also been discussed above. Injector subsystem **15a** can send information about its state or its programs to imaging subsystem **15b** such as, but not restricted to:

- Contrast enhancing fluid type or volume, read by bar code or operator input
- Contrast enhancing fluid concentration
- Time since mixing of the contrast enhancing fluid
- Time since last injection
- Time since start of current injection
- Time of last servicing
- Injector system unique identification number or code
- Injector hardware or software configuration (version number)
- Status of agent agitation, for instance, mixing or ready
- Current Selected Infuse Rate
- Current Selected Bolus Rate
- Current Selected Bolus Volume
- Current injection Phase
- Actual Flow Rate
- Actual Volume Delivered Total Injected Volume

Volume Remaining
 Bolus Ready
 Bolus Low Volume
 Bolus Running
 Bolus Complete
 Infuse Ready
 Infuse Running
 Infuse Pending
 Injection Stopped or Canceled
 Injector Error Detected and Type

Information such as set forth above can be used by imager subsystem **15b** to adjust its operation to optimize the image acquisition process as described above. It can also be displayed on the imager subsystem operator interface so that it is convenient for the operator to assess the injector subsystem's functioning. It may be that imager subsystem **15b** cannot automatically adjust its operation based upon the transmitted information, but that the operator has to make an adjustment. The information could be displayed at the imager to allow the operator to view the information before making adjustments. This may be for regulatory approval or to simplify subsystem design verification.

Information can also be transmitted from imager subsystem **15b** to injector subsystem **15a**. This information may be used to adjust the injector subsystem operation as mentioned above, or it may simply be displayed so the operator at the injector user interface **150a** can be informed about key imager subsystem information. A basic benefit is the ability to start and stop the injection from imager operator interface **150b**. A much more sophisticated capability is to be able to program all injector functions from the imager subsystem operator interface **150b** as if the operator were at injector subsystem operator interface **150a**. An example of a middle ground is if injector subsystem **15a** communicates possible preprogramming sequences to imager subsystem **15b**, and the operator at imager operator interface **150b** chooses among the sequences.

Another benefit of data communication between subsystems is the ability for imager subsystem **15b** or the operator at imager subsystem operator interface **150b** to determine that an image or measurement was not sufficient and command the injector to repeat the injections or to perform a different injection. A planned example of this is what is often termed a test injection. After an injection is made and a series of images or measurements are acquired, scanner/imager processor unit **92b** or injector processor unit **92a** can determine from the information gathered the injection parameters, including timing, that are most likely to create the desired enhancement in the final image or measurement.

Given the criticality of the information being transmitted between the two subsystems, it is desirable that there be significant safeguards to prevent error or problems. In a preferred embodiment, the electrical signals of the two subsystems are not electrically connected to each other, which can be accomplished in many ways. Optical isolators, isolation transformers and capacitance based isolators commonly used components for providing electrical isolation. Other means of isolation are by the use of transmitted or broadcast electromagnetic or ultrasonic energy for the transmission means.

It is also preferable that each subsystem be able to operate even if interface **90c** is not operating. This result can be accomplished within the programming of each subsystem processor unit. If the operator tries to use a function that requires communications between subsystems, and commu-

nications are not available or in an unknown state, the operator is preferably either informed of that condition, or an attempt to communicate is made and the operator is informed of the results of that attempt.

It is also preferable to structure data transmission so that errors are avoided or tolerated. The discussion below provides an example procedure to provide error detection and tolerance. The example discusses information transmission from injector subsystem **15a** to imager subsystem **15b** with acknowledgement of proper receipt by imager subsystem **15b** being sent to injector subsystem **15a**, but the same methodology can be readily applied with the respective roles reversed to transmit information in the other direction. The use of two separate transmission channels can be used as discussed below. A single transmission channel can also be used.

EXAMPLE

In one embodiment, injector subsystem **15a** periodically transmits status information through subsystem interface **90c** to imager subsystem **15b**, which preferably only has to acknowledge the successful (or unsuccessful) data reception. In this embodiment, imager subsystem **15b** does not request data from injector subsystem **15a** or send data other than acknowledgements to injector subsystem **15a**. A message based flow control mechanism is preferably used to regulate the data flow from injector subsystem **15a**.

In this embodiment, only two different messages are preferably sent from injector subsystem **15a** to imager subsystem **15b**. The injector subsystem interface status message preferably contains the interface software version number and indicates if injector status information is available, and the injector status message preferably contains, for example, injection parameter values and volume information. The injector subsystem interface status message is preferably transmitted when connection is established or resumed, which is determined by the status of the handshake lines. It is preferably sent approximately every 500 milliseconds (not including retries) as long as injector status information is not yet or no longer available. If injector status information is available and the injector subsystem interface status message has been acknowledged, the injector status message is preferably transmitted approximately every 500 milliseconds (not including retries).

The following discussion provides a detailed description of the communication protocol and the format of the two messages discussed above.

Data Exchange Mode

Communication may, for example, be serial, asynchronous and full duplex at 19200 Baud with flow control using RTS/CTS. Each data frame sent between the two subsystems preferably comprises ten bits: one start bit (logic 0), eight data bits (LSB first, no parity), and one stop bit (logic 1). The start and stop bits preferably have the same width as data bits. The electrical interface is preferably designed to be compatible with the EIA RS-232.V28 standard.

Message Format

Messages sent from injector subsystem **15a** to imager subsystem **15b** preferably have the format set forth in Table 1:

TABLE 1

STX	Sequence Number	Message Length	Message Data	CRC	ETX
1 byte	1 (2) byte	1 (2) byte	up to 48 (96) byte	1 (2) byte	1 byte

Numbers in parenthesis in Table 1 represent the actual size during transmission after data folding has been performed. Data folding is discussed below.

STX

The STX (start of text) character indicates the beginning of a message. The one byte hexadecimal code for STX is, for example, 0x02.

Sequence Number

A one byte sequence number is preferably used to enable the receiving system to detect a retransmitted message. The sequence number is necessary for the case that a message is successfully transmitted and received but the receiver's acknowledge becomes corrupted. The transmitting system preferably retransmits the message after a response time-out has occurred. As the receiver already received the message successfully the first time, it can identify the message as a duplicate by checking the sequence number against the sequence number of the last received message. The receiver preferably acknowledges and then discards the duplicate message. The receiver, therefore, preferably maintains a receive sequence number which is set to the sequence number of a received message after it has been validated. The transmitter also preferably maintains a transmit sequence number which represents the sequence number of the last message for which a positive acknowledge was received. An example of internal transmit and receive sequence numbering for a message transmission and the receiver's response is illustrated in FIG. 4.

Message Length

The one byte Message Length field specifies the size of the following Message Data field in bytes. The Message Length value range is, for example, 1 to 48.

Message Data

This portion of the message is the data portion of the message. If a message contains data items comprising multiple bytes, the byte order is preferably little endian (least significant byte first). Examples of supported messages and associated message format are discussed in further detail below.

CRC

An eight bit CRC is preferably used to secure the data to be transmitted. It enables the receiving system to detect message data corruption. The CRC covers the Sequence Number, Message Length, and Message Data fields before folding. An example of a polynomial used to calculate the CRC is: $x^8+x^4+x^2+x+1$.

ETX

The ETX (end of text) character indicates the end of a message. The one byte hexadecimal code for ETX is, for example, 0x03.

Message Acknowledge

As mentioned above, the receiving system preferably acknowledges each received message by sending a single control character. The transmitting system preferably does not transmit a new message until the last transmitted message has been acknowledged or a response time-out has occurred. An acknowledge can be positive (ACK), indicating successful message reception and CRC validation, or negative (NAK), indicating an invalid CRC or a receive

time-out. The injector preferably retransmits a message until it receives a positive acknowledge. If a status message has to be retransmitted and updated status information is available, a new message with the updated status is preferably sent instead. The one byte hexadecimal code for an ACK is, for example, 0x06. A NAK is represented as hexadecimal 0x15.

Data Folding

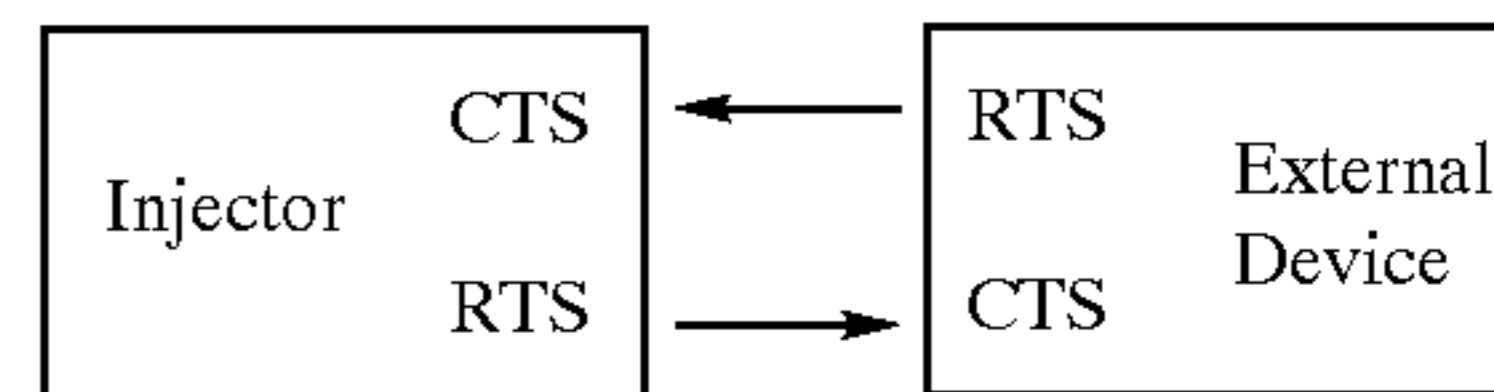
To avoid data escaping, all bytes between the STX and ETX control characters are preferably 'folded' to guarantee that their values fall within the printable ASCII character range (0x20 to 0xE7) and do not overlap with control characters. This result is achieved by splitting each byte into two nibbles and adding a fixed offset to each nibble. The offset to be used is preferably 0x30. Folding the data between the STX and ETX, therefore, doubles the size of a message. An example message and the folded data to be transmitted are given below in Table 2:

TABLE 2

	Original Message	Folded Message
STX	0x02	0x02
Sequence Number	0x10	0x30, 0x31
Message Length	0x02	0x32, 0x30
Message Data	0x45, 0x67	0x35, 0x34, 0x37, 0x36
CRC	0xB6	0x36, 0x3B
ETX	0x03	0x03

Handshaking

Two hardware signals (CTS and RTS) may be used to implement a message based flow control as described above. The signals may also be used to detect connection between injector subsystem 15a and imager subsystem 15b. FIG. 5 illustrates the signals and how they are connected.



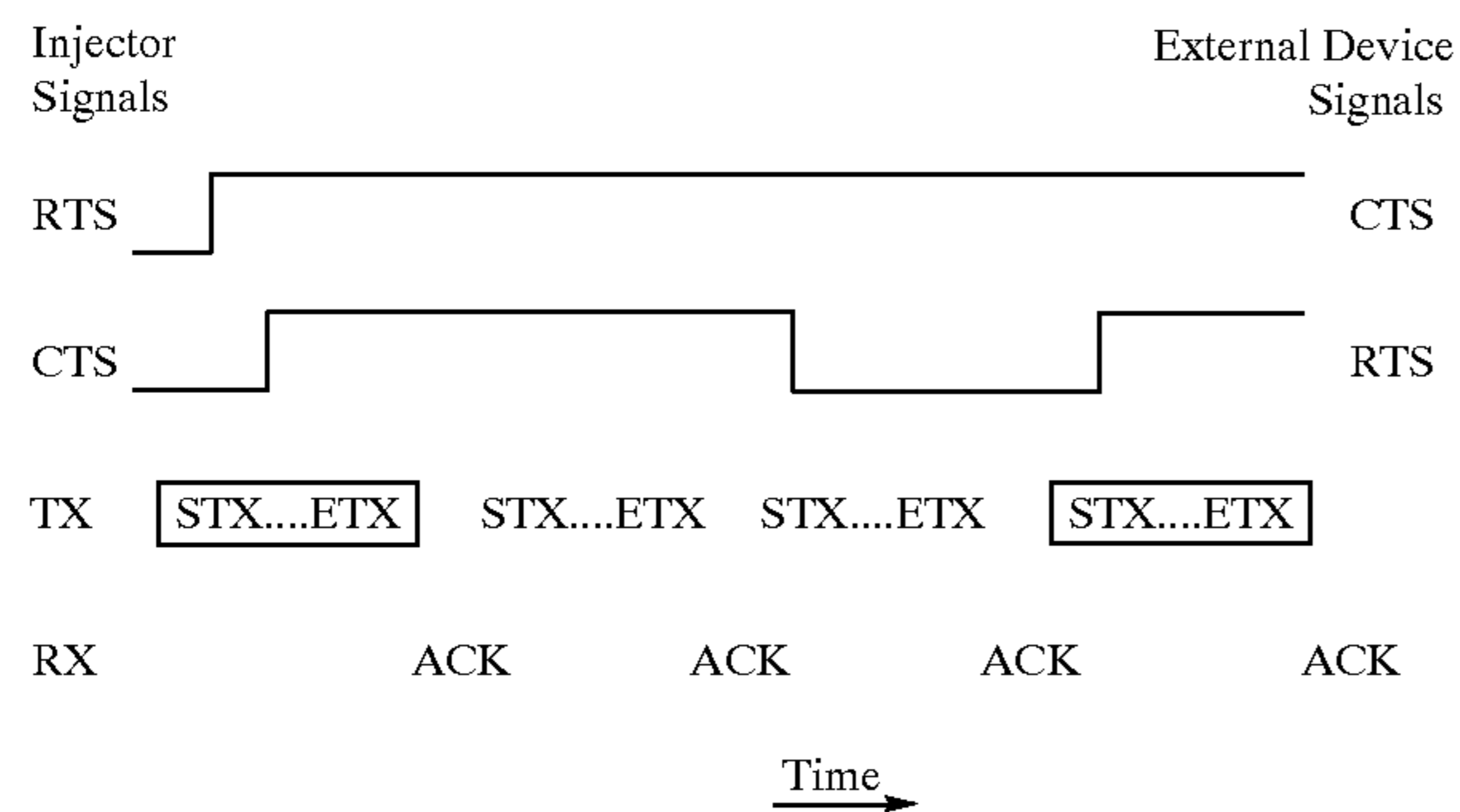
Connection Establishment

The injector preferably activates RTS once it is ready to transmit data to imager subsystem 15b. The RTS signal preferably stays active as long as injector subsystem 15a is powered up and operating normally. Initial connection is preferably established when injector subsystem 15a has activated its RTS output and detects its CTS input as active for the first time. Connection is preferably resumed when injector subsystem 15a detects its CTS input as active after it had been deactivated. Imager subsystem 15b can activate its RTS output (the injector's CTS input) at any time; it is not required to wait for injector subsystem 15a to activate its RTS output first.

Flow Control

Injector subsystem 15a preferably transmits the injector subsystem interface status message when connection is initially detected or resumed. Once injector status information is available and the injector subsystem interface status message has been acknowledged (ACK), injector subsystem 15a preferably transmits the injector status message about every 500 milliseconds, not including retransmits. As long as imager subsystem 15b does not deactivate its RTS signal, injector subsystem 15a preferably continues its periodic transmission. If imager subsystem 15b deactivates its RTS output, injector subsystem 15a preferably finishes the cur-

rent message transmission. It then waits for imager subsystem 15b to activate its RTS signal again. When the imager subsystem's RTS output is active again, injector subsystem 15a preferably restarts transmission with an injector subsystem interface status message. FIG. 6 illustrates a flow control example: (INJECTOR SUBSYSTEM interface status messages are indicated by a solid border)



Time-Outs

The requirement that each successfully received message be acknowledged by the receiver, has the side effect that the transmitter can potentially wait forever for an acknowledge (ACK/NAK). Another possible error condition is that the transmitter aborts the transmission before the ETX character is sent. A set of time-outs are preferably used to allow the transmitting and receiving systems to recover from these situations.

A time-out duration may, for example, be based on the following calculations: At a baud rate of 19200 it takes about 52 microseconds to transfer one data bit. One data byte (a 10 bit frame) can, therefore, be transmitted in about 520 microseconds. Assuming no delay between bytes, transmission of a maximum length message (104 bytes, folded) takes approximately 54.1 milliseconds. The required one byte response from the receiver adds another 520 microseconds, not taking any validation and processing of the received message into account.

Transmit Time-out

A message transmission time-out may be used to account for situations in which the transmission of a message never completes or takes too long because of an internal transmitter problem. If a transmission time-out is not incorporated, the injector could possibly wait forever for the completion of a message transmission which would block the transmission of all subsequent messages. As previously discussed, a maximum length message can be transmitted in a minimum of 54.1 milliseconds. Allowing some delay between bytes because of processing overhead, the transmit time-out may, for example, be set to 60 milliseconds. The transmit time-out period begins with the STX character being transmitted.

Receive Time-out

A message receive time-out may be used to account for situations in which the reception of a message never completes (e.g. ETX missing) or takes too long. If a receive time-out is not incorporated into the design the imager subsystem could possibly wait forever for the reception of a message to complete which would block the reception of all subsequent messages. Since the timing associated with the reception of a message is dictated by the transmitter, the same value as for the transmit time-out (60 milliseconds) may be used. The receive time-out period begins when the STX character is received.

Response Time-out

A message response time-out may be used to account for situations in which imager subsystem 15b does not acknowledge or takes too long to acknowledge a received message. Even though imager subsystem 15b does not send any messages in this embodiment and an acknowledge could therefore be expected after a few milliseconds, the response time-out is also preferably set to 60 milliseconds. The response time-out period preferably begins when the ETX character is transmitted.

The section below provides a brief summary of the handling of a number of error situations by the present communication interface system.

Error Handling

Injector Subsystem

Transmission Time-out: The time to transmit an entire message exceeded the transmission time-out period. This error indicates an internal problem of the injector transmitter. The injector will deactivate its RTS signal, indicating that no further messages will be sent to the imager subsystem. RTS will remain deactivated until power is cycled off/on and the injector is ready to transmit data again.

Response Time-out: The imager subsystem did not respond to a message within the response time-out period. The injector will retransmit the last message unless updated status information is available which would be transmitted instead of the last message.

NAK Received: The imager subsystem did not properly receive a message (CRC mismatch, receive time-out). The injector will retransmit the last message unless updated status information is available which would be transmitted instead of the last message.

Imager Subsystem

Receive Time-out: The time that it took to receive a message exceeded the receive time-out period. The imager subsystem will send a NAK to request a retransmit of the message.

STX Character Missing: Other data are received when waiting for an STX character. The imager subsystem will discard all characters until an STX is received. If the injector was trying to send a message, a response time-out will occur and the message will be sent again.

CRC Mismatch: The CRC calculated from the received data did not match the received CRC value. The imager subsystem will send a NAK to the injector after the receive time-out period has expired.

Duplicate Message Number: The received message number was equal to the last message number received. The imager subsystem will send an acknowledge and disregard the retransmitted message.

Dropped Data Bytes: During reception of a message one or more data bytes were lost. This error will most likely result in a CRC mismatch. The imager subsystem will send a NAK to the injector after the receive time-out period has expired.

Message Data Format

The Message Data portion of a message as discussed above may have the format set forth in Table 3 below.

TABLE 3

ID	ID specific data
(1 byte)	(up to 47 bytes)

Supported IDs

Examples of supported messages and associated data for external device interface status and injector status are pro-

vided in Tables 4 and 5, respectively. Binary data of word size are transmitted in little endian byte order. FIG. 7 illustrates a state transition diagram of the receive processing to be performed by an external device (for example, imager subsystem 15b) communicating with injector subsystem 15a.

TABLE 4

Field	Encoding	Data Type	Value Range	Description
ID Data	Binary	Byte 7	0xF5	
Injector Status Data Available	Binary	Bit 7	0 or 1	The bit is 1 when injector status information is available for transmission.
Reserved	N/A	Bit 6 - 0	N/A	
SW Version	ASCII	Char [4]	'1.0' to '99.9'	
Reserved	N/A	Byte [2]	N/A	

TABLE 5

Field	Encoding	Data Type	Value Range	Description
ID Data	Binary	Byte	0xFA	
Programmed Infusion Rate	Binary	Word	0 to 290	The value is given in tenths ml/min. The actual range therefore is 0.0 to 29 ml/min. Resolution: 0.5 ml/min between 0.0 and 10, 1 ml/min between 10 and 29
Programmed Bolus Rate	Binary	Word	0 to 30	The value is given in tenths ml/s. The actual range therefore is 0.0 to 3.0 ml/s. Resolution: 1.0 ml/s
Programmed Bolus Volume	Binary	Word	0 to 30	The value is given in ml. Resolution: 1 ml
Flow Rate	Binary	Word	0 to 1800	The value represents a non-cumulative running average while an injection is in progress. It will be 0 before and after an injection. The value is given in tenths ml/min. The actual range therefore is 0.0 to 180 ml/min. During a bolus, the external device should convert the received flow rate value to units of ml/s.
Total Injected Volume	Binary	Word	0 to 9990	The value is given in tenths ml. The actual range therefore is 0.0 to 999 ml. Resolution: 0.5 ml between 0.0 and 100, 1 ml between 100 and 999
Volume Remaining	Binary	Word	0 to 300	The value is given in tenths ml. The actual range therefore is 0.0 to 30.0 ml. Resolution: 0.5 ml
Bolus Ready	Binary	Bit 7	0 or 1	The bit is 1 when the injector is ready for an injection and both bolus parameters are >0. The bit is 0 at any other time, including during a bolus.
Infusion Ready	Binary	Bit 6	0 or 1	The bit is 1 when the injector is ready for an injection and the infusion rate is >0. The bit is 0 at any other time, including during an infusion

TABLE 5-continued

Field	Encoding	Data Type	Value Range	Description
Bolus Running	Binary	Bit 5	0 or 1	The bit is 1 while the injector is performing a bolus. The bit is 0 at any other time.
Infusion Running	Binary	Bit 4	0 or 1	The bit is 1 while the injector is performing an infusion. The bit is 0 at any other time.
Low Volume	Binary	Bit 3	0 or 1	The bit is 1 if the programmed bolus volume is less than the volume remaining. If this condition becomes true during a bolus, Low Volume will not be indicated until the bolus ends to avoid operator confusion. The bit is 0 at any other time.
Infusion Pending	Binary	Bit 2	0 or 1	The bit is 1 if an infusion was interrupted by or requested during a bolus and will resume after the bolus ends. The bit is 0 at any other time.
Bolus Complete	Binary	Bit 1	0 or 1	The bit is 1 when a bolus completes (programmed volume is delivered). It is 0 at any other time. Note that the bit is only set for one message. The external device is responsible for latching this information if desired.
Injection aborted/canceled	Binary	Bit 0	0 or 1	The bit is 1 when an injection is aborted or canceled e.g. due to an invalid key press or running out of volume. The bit is 0 at any other time. Note that the bit is only for one status message. The external device is responsible for latching this information if desired.
Error Code	ASCII	Char [3]	' ' or 'A 1' to 'Z99'	Three spaces indicate no error. The currently supported error codes are listed below: P1: Perform check for air P2: No volume remaining P3: Pressure stall P4: Syringe attached P5: Syringe removed S1: Disarm due to key press C: Critical Error 'P' codes require an operator acknowledge, 'S' codes time out after five seconds 'C' requires injector power to be cycled.

Layering

A summary of a preferred embodiment of the layering involved in the present example is presented in Table 6.

TABLE 6

OSI Layer	Key Layer Functions/Services	Preferred Methods	Other Representative Methods
Layer 1. — Physical Layer	Physical connections, physical service data units (n-bit transmission), physical connection end points, sequencing, data circuit identification, fault condition notification, quality of service parameters (error rates, service availability, transmission rates, delays)	RS-232.V28	thinnet, thicknet, unshielded twisted pair (UTP), RS-232-C, FDDI, ISN, wireless RF, wireless infrared
Layer 2. — Data Link Layer	Data link connection, data link service data units, error notification, flow control, quality of service parameters	Ack>/<Nak> message control, start of message <Stx> and end of message <Etx> formatting, message numbering scheme, transmit and receive retries (optional), duplicate message number detection and handling, CRC-16 error detection and correction operations	Ethernet, token ring, HDLC
Layer 3. — Network Layer	Data packet routing	Not used	Internet Protocol (IP, SLIP, CSLIP, PPP), Internet Control Message Protocol (ICMP), Internet Group Management Protocol (IGMP)
Layer 4. — Transport Layer	Error recovery and flow control, message multiplexing	Not used	Transmission Control Protocol (TCP), User Datagram Protocol (UDP)
Layer 5. — Session Layer	Control structure for managing communications, session connection, session management, session termination, interaction management	Outgoing data is not sent (session not initiated) unless external device enables communication through CTS	Remote Procedure Call (RPC)
Layer 6. — Presentation Layer	Common operations on the structure of the data being exchanged, syntax conversion, syntax negotiation, encryption, compression	Message coding and decoding, data folding	External Data Representation (XDR)
Layer 7. — Application Layer	Communication management between applications, synchronization of applications, availability determination, file transfer access, job transfer	Fixed protocol, no negotiation. Injector and imager share timing and control information to perform the imaging procedure.	SMTP, SNMP, ftp, telnet, DNS, INS, NFS, arp, rlogin, talk, ntp, traceroute

All medical products are regulated by government bodies in some way. This approval process includes validation and verification that the equipment performs properly under all conditions. As discussed above, it is possible that injector subsystem **15a** will be designed and manufactured by one company and that imager subsystem **15b** will be designed and manufactured by another. In fact, there may be multiple manufacturers for each. Subsystem interface **90c** may be manufactured by any of the above, or even be manufactured by a totally different company. As also discussed above, the equipment functions may be coordinated by a human operator to perform the imaging needed to determine the diagnosis. It is very difficult to test a piece of equipment under all possible conditions when it has to communicate and cooperate with another piece of equipment. Even if two pieces of equipment are manufactured by the same company, they are often designed by separate teams and there are thus still coordination and testing difficulties. In addition, there may

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be a problem with changes or improvements to one piece of equipment being compatible with older or newer versions of the other equipment. It is, therefore, desirable that standard interfaces, such as the EIA RS-232.V28 standard be used in the present invention. To sufficiently isolate injector and imager subsystems **15A** and **15a**, it is preferred that subsystem interface **90c** be powered separately and be isolated from both of the subsystems. This has a regulatory and validation benefit.

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Although the present invention has been described in detail in connection with the above embodiments and/or examples, it is to be understood that such detail is solely for that purpose. For example, many of the functions separately called out and described can be performed by identical pieces of hardware. These and other variations can be made by those skilled in the art without departing from the spirit of the invention except as it may be limited by the following claims.

What is claimed is:

1. A system for administering an active substance to a patient comprising:
 - a source of a contrast or enhancement medium;
 - at least a first delivery unit in fluid connection with the source of contrast or enhancement medium;
 - at least a second delivery unit in fluid communication with an active substance;
 - an imaging energy source adapted to apply imaging energy to a region of the patient;
 - an imaging unit providing an indication of an internal view of the patient based upon a signal resulting from the imaging energy applied to the region of the patient, the signal being affected by a condition of the contrast or enhancement medium in the patient; and
 - a control unit adapted to control the administration of the active substance based on the indication.
2. The system of claim 1 wherein the active substance comprises at least one of a therapeutic drug and a gene therapy.
3. The system of claim 1 wherein the active substance comprises a diagnostic substance.
4. The system of claim 1 wherein the energy applied to the region of the patient is sonic energy and the signal is an acoustical signal.
5. The system of claim 1 wherein the energy source is further operable to activate the active substance.
6. The system of claim 1 wherein the control unit is further adapted to control the at least a second delivery unit.
7. The system of claim 1 wherein the control is further adapted to control the at least the first and the at least the second delivery unit.
8. The system of claim 1, further comprising:
 - an activating energy source adapted to activate the active substance.
9. The system of claim 8 wherein the imaging energy source and the activation energy source provide the same type of energy.
10. The system of claim 8 wherein the imaging energy source and the activation energy source is sonic energy.
11. The system of claim 1, further comprising:
 - a source of the active substance.
12. The system of claim 1 wherein the source of contrast or enhancement medium further comprises the active substance.
13. A system for administering an active substance to a patient comprising:
 - a source of a contrast or enhancement medium;
 - at least one delivery unit in fluid communication with both the source of contrast or enhancement medium and an active substance;
 - an imaging energy source adapted to apply imaging energy to a region of the patient;
 - an imaging unit providing an indication of an internal view of the patient based upon a signal resulting from the imaging energy applied to the region of the patient, the signal being affected by a condition of the contrast or enhancement medium in the patient; and
 - a control unit adapted to control the administration of the active substance based on the indication.
14. The system of claim 13 wherein the active substance comprises at least one of a therapeutic drug and a gene therapy.
15. The system of claim 13 wherein the active substance comprises a diagnostic substance.

16. The system of claim 13 wherein the energy applied to the region of the patient is sonic energy and the signal is an acoustical signal.
17. The system of claim 13 wherein the energy source is further operable to activate the active substance.
18. The system of claim 13 wherein the control unit is further adapted to control the at least one delivery unit.
19. The system of claim 13, further comprising:
 - an activating energy source adapted to activate the active substance.
20. The system of claim 19 wherein the imaging energy source and the activation energy source provide the same type of energy.
21. The system of claim 19 wherein the imaging energy source and the activation energy source is sonic energy.
22. The system of claim 13, further comprising:
 - a source of the active substance.
23. The system of claim 13 wherein the source of contrast or enhancement medium further comprises the active substance.
24. A system for administering an active substance to a patient comprising:
 - a source of a contrast or enhancement medium;
 - at least one delivery unit in fluid communication with one or both of the source of contrast or enhancement medium and an active substance for delivering the contrast or enhancement medium or the active substance to the patient, wherein the active substance comprises at least one of a therapeutic drug and a gene therapy;
 - an imaging energy source adapted to apply imaging energy to a region of the patient;
 - an imaging unit providing an indication of an internal view of the patient based upon a signal resulting from the imaging energy applied to the region of the patient, the signal being affected by a condition of the contrast or enhancement medium in the patient; and
 - a control unit adapted to control the administration of the active substance based on the indication.
25. The system of claim 24 wherein the energy applied to the region of the patient is sonic energy and the signal is an acoustical signal.
26. The system of claim 24 wherein the energy source is further operable to activate the active substance.
27. The system of claim 24 wherein the control unit is further adapted to control the at least one delivery unit.
28. The system of claim 24, further comprising:
 - an activating energy source adapted to activate the active substance.
29. The system of claim 28 wherein the imaging energy source and the activation energy source provide the same type of energy.
30. The system of claim 28 wherein the imaging energy source and the activation energy source is sonic energy.
31. The system of claim 24, further comprising:
 - a source of the active substance.
32. The system of claim 24 wherein the source of contrast or enhancement medium further comprises the active substance.
33. The system of claim 24 wherein the at least one delivery unit comprises at least two delivery units, a first delivery unit in fluid communication with the source of contrast or enhancement medium and a second delivery unit in fluid communication with the active substance.
34. A method for administering an active substance to a patient comprising:

providing a source of a contrast or enhancement medium;
 providing at least one delivery unit in fluid connection
 with one or both of the source of contrast or enhance-
 ment medium and an active substance for delivering the
 contrast or enhancement medium or the active sub-
 stance to the patient, wherein the active substance
 comprises at least one of a therapeutic drug and a gene
 therapy;
 injecting the contrast or enhancement medium into the
 patient;
 providing an imaging energy source adapted to apply
 imaging energy to a region of the patient;
 applying energy to the region of the patient;
 providing an imaging unit providing an indication of an
 internal view of the patient based upon a signal result-
 ing from the imaging energy applied to the region of the
 patient, the signal being affected by a condition of the
 contrast or enhancement medium in the patient;
 generating an indication of an internal view of the patient;
 and
 controlling the administration of the active substance
 based on the indication.

35. The method of claim **34** wherein the energy applied to
 the region of the patient is sonic energy and the signal is an
 acoustical signal.

36. The method of claim **34** wherein the energy source is
 further operable to activate the active substance.

37. The method of claim **34** wherein the control unit is
 further adapted to control the at least one delivery unit.

38. The method of claim **34**, further comprising:
 providing an activating energy source adapted to activate
 the active substance.

39. The method of claim **38** wherein the imaging energy
 source and the activation energy source provide the same
 type of energy.

40. The method of claim **38** wherein the imaging energy
 source and the activation energy source is sonic energy.

41. The system of claim **34**, further comprising:
 a source of the active substance.

42. The system of claim **34** wherein the source of contrast
 or enhancement medium further comprises the active sub-
 stance.

43. A system for administering an active substance to a
 patient comprising:
 a source of an ultrasound contrast or enhancement
 medium containing the active substance;
 at least one delivery unit in fluid communication with the
 source of contrast or enhancement medium and an
 active substance for delivering the contrast or enhance-
 ment medium or the active substance to the patient;
 an imaging energy source adapted to apply imaging
 energy to a region of the patient;
 an imaging unit providing an indication of an internal
 view of the patient based upon a signal resulting from
 the imaging energy applied to the region of the patient,
 the signal being affected by a condition of the contrast
 or enhancement medium in the patient; and
 a control unit adapted to control the administration of the
 active substance based on the indication.

44. The system of claim **43** wherein the energy applied to
 the region of the patient is sonic energy and the signal is an
 acoustical signal.

45. The system of claim **43** wherein the energy source is
 further operable to activate the active substance.

46. The system of claim **43** wherein the control unit is
 further adapted to control the at least one delivery unit.

47. The system of claim **43**, further comprising:
 an activating energy source adapted to activate the active
 substance.

48. The system of claim **43** wherein the imaging energy
 source and the activation energy source provide the same
 type of energy.

49. The system of claim **47** wherein the imaging energy
 source and the activation energy source is sonic energy.

50. A system for administering an active substance to a
 patient comprising:
 a source of a contrast or enhancement medium;
 at least one delivery unit in fluid communication with one
 or both of the source of contrast or enhancement
 medium and the active substance for delivering the
 contrast or enhancement medium or an active substance
 to the patient, wherein the active substance comprises
 at least one of a therapeutic drug and a gene therapy;
 an imaging energy source adapted to apply imaging
 energy to a region of the patient and to activate the
 active substance;
 an imaging unit providing an indication of an internal
 view of the patient based upon a signal resulting from
 the imaging energy applied to the region of the patient,
 the signal being affected by a condition of the contrast
 or enhancement medium in the patient; and
 a control unit adapted to control the administration of the
 active substance based on the indication.

51. A system for administering an active substance to a
 patient comprising:
 a source of a contrast or enhancement medium;
 at least one delivery unit in fluid communication with one
 or both of the source of contrast or enhancement
 medium and the active substance for delivering the
 contrast or enhancement medium or an active substance
 to the patient, wherein the active substance comprises
 at least one of a therapeutic drug and a gene therapy;
 an imaging energy source adapted to apply imaging
 energy to a region of the patient;
 an imaging unit providing an indication of an internal
 view of the patient based upon a signal resulting from
 the imaging energy applied to the region of the patient,
 the signal being affected by a condition of the contrast
 or enhancement medium in the patient; and
 a control unit adapted to control the at least one delivery
 unit and the administration of the active substance
 based on the indication.

52. A system for administering an active substance to a
 patient comprising:
 a source of a contrast or enhancement medium;
 at least one delivery unit in fluid communication with one
 or both of the source of contrast or enhancement
 medium and the active substance for delivering the
 contrast or enhancement medium or an active substance
 to the patient, wherein the active substance comprises
 at least one of a therapeutic drug and a gene therapy;
 an imaging energy source adapted to apply imaging
 energy to a region of the patient;
 an activating energy source adapted to activate the active
 substance;
 an imaging unit providing an indication of an internal
 view of the patient based upon a signal resulting from
 the imaging energy applied to the region of the patient,
 the signal being affected by a condition of the contrast
 or enhancement medium in the patient; and

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a control unit adapted to control the administration of the active substance based on the indication.

53. A system for administering an active substance to a patient comprising:

a source of a contrast or enhancement medium;

at least one delivery unit in fluid communication with one or both of the source of contrast or enhancement medium and an active substance for delivering the contrast or enhancement medium or the active substance to the patient, wherein the active substance comprises at least one of a therapeutic drug and a gene therapy;

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a sonic imaging energy source adapted to apply imaging energy to a region of the patient;

a sonic activating energy source adapted to activate the active substance;

5 an imaging unit providing an indication of an internal view of the patient based upon a signal resulting from the imaging energy applied to the region of the patient, the signal being affected by a condition of the contrast or enhancement medium in the patient; and

10 a control unit adapted to control the administration of the active substance based on the indication.

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